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VAPORIZATION RESPONSE OF EVAPORATING DROPS WITH FINITE THERMAL CONDUCTIVITY

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16. Abstract

A numerical computing procedure was developed for calculating vaporization histories of evaporating drops in a combustor in which travelling transverse oscillations occurred. The liquid drop was assumed to have a finite thermal conductivity. The system of equations was solved by using a finite difference method programmed for solution on a high speed digital computer. Oscillations in the ratio of vaporization of an array of repetitively injected drops in the combustor were obtained from summation of individual drop histories. A nonlinear in-phase frequency response factor for the entire vaporization process to oscillations in pressure was evaluated. In addition, a nonlinear out-of-phase response factor, in-phase and out-of-phase harmonic response factors, and a Princeton type "n" and "τ" were determined. The resulting data was correlated and is presented in graphical format. Qualitative agreement with the open literature is obtained in the behavior of the in-phase response factor. Quantitatively, however, the results of the present finite conductivity spray analysis do not correlate with the results of a single drop model.

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VAPORIZATION RESPONSE OF EVAPORATING DROPS WITH FINITE THERMAL CONDUCTIVITY

by V. D. Agosta and S. S. Hammer

SUMMARY

The primary objective of the analysis was to obtain a numerical computing procedure for determining the vaporization response of droplets with finite thermal conductivity in an oscillating pressure and flow field. The governing equations for vaporization of liquid drops in a rocket combustor environment were taken from Refs. 1 and 2. Additional equations were employed to account for finite thermal conductivity of the liquid drop (Ref. 3). The system of equations were solved utilizing a finite difference technique and a high speed digital computer. Oscillation in the rates of vaporization of an array of repetitively injected drops in the combustor were obtained from the summation of individual drop histories. A nonlinear in-phase frequency response of the entire vaporization process to pressure oscillations was calculated and a response factor, $R_{n\ell}$, was determined as defined by Equation 1 of Ref. 4. In addition, a nonlinear out-of-phase response factor, $I_{n\ell}$ in-phase and out-of-phase harmonic response factors R_1, R_2, I_1, I_2 and a Princeton type "n" and "τ" were determined as described in Refs. 5 and 6. In general, it was found that the nonlinear in-phase response factor, $R_{n\ell}$, was not very sensitive to variations of up to 10^3 in the liquid thermal conductivity, for the frequency range of interest in combustion instability studies.

The resulting data was correlated and is presented in graphical format. Qualitative agreement with the open literature is obtained in the behavior of the in-phase response factor.

INTRODUCTION

Studies on nonlinear combustion instability have been performed (Refs. 2,4,7) which indicate that vaporizing drops are sensitive to frequency-dependent pressure oscillations. The sensitivity of the vaporization processes has been traced to thermal time lags, namely, that time delay between changes in the drop temperature and subsequent mass vaporization and changes in the drop environment. The thermal time lag model proposed in Ref. 7 is extended to include the effects of drop thermal conductivity on drop surface temperature and mass vaporization.

Wave deformation effects on droplet vaporization as proposed in Ref. 5 were also considered in this investigation by varying the harmonic distortion in a pressure disturbance propagating in a liquid rocket combustion chamber.

Nonlinear and harmonic in-phase and out-of-phase response factors which have been evolved from linear limit-cycle stability analyses have been adopted for use in this report. This procedure is consistent with present usage and allows for the comparison of data and results for similar parameters and factors.

THEORY

Drops of liquid oxygen are assumed to be vaporizing in combustion gases, composed of stoichiometric reaction products with hydrogen in a cylindrical combustion chamber, with an established travelling-transverse acoustic mode. The instantaneous pressure and gas velocity functions consist of the steady state and oscillating components attributed to the acoustic mode. They are expressed as

$$p = \bar{p}_c \left(1 + .859 J_1 \left(\frac{1.841 R}{R_w}\right)\right) \left(\frac{\Delta p_1}{\bar{p}_c} \cos(2\pi ft - \theta) + \frac{\Delta p_2}{\bar{p}_c} \cos(4\pi ft - \varphi)\right) \qquad (1)$$

$$U = \left[U_a^2 + U_R^2 + U_\theta^2\right]^{\frac{1}{2}} \qquad (2)$$

$$U_R = .430 \frac{a}{\gamma}\left(J_o\left(\frac{1.841R}{R_w}\right) - J_2\left(\frac{1.841R}{R_w}\right)\right)\left(\frac{\Delta p_1}{\bar{p}_c}\sin(2\pi ft-\theta) + \frac{\Delta p_2}{\bar{p}_c}\sin(4\pi ft-\varphi)\right) \quad (3)$$

$$U_\theta = .467 \frac{a}{\gamma}\frac{R_w}{R} J_1\left(\frac{1.841R}{R_w}\right)\left(\frac{\Delta p_1}{\bar{p}_c}\cos(2\pi ft-\theta) + \frac{\Delta p_2}{\bar{p}_c}\cos(4\pi ft-\varphi)\right) \quad (4)$$

$$U_a = U_{a_f}\left(1 - \frac{m}{m_i}\right. \quad (5)$$

where

a = speed of sound
f = frequency of oscillations
J = Bessel function
m = mass of droplet
m_i = initial mass of droplet
p = instantaneous pressure
$\bar{p}_c$ = mean chamber pressure
Δp_1 = peak-to-peak pressure amplitude (fundamental)
Δp_2 = peak-to-peak pressure amplitude (harmonic)
R = radial location in chamber
R_N = radius of chamber
t = time
T = temperature
U = gas velocity
U_{a_f} = final axial gas velocity
θ = phase angle
γ = isentropic exponent

The wave is assumed to be adiabatic with chamber temperature and pressure related by

$$\frac{T_c}{\bar{T}_c} = \left(\frac{p}{\bar{p}_c}\right)^{\frac{\gamma-1}{\gamma}} \quad (6)$$

The oscillations of pressure and velocity are superimposed on the mean level of parameters affecting drop evaporation and motion. The vaporization rate is controlled by heat and mass transfer to the surface of a drop with finite thermal conductivity. Drop motion is controlled by a momentum balance as a result of drag with the enveloping gas. Axial gas velocity is proportional to the fraction of drop mass vaporized, and it attains a final assumed velocity at complete evaporation. The complete drop history is defined by the following equations for weight evaporation rate heat transfer, acceleration in an axial direction and temperature distribution within the drop.

The governing differential equation for temperature in a spherical liquid droplet is given by

$$T_t = \alpha\left(T_{rr} + \frac{2}{r} T_r\right) \qquad \begin{matrix} 0 \leq r \leq r_s \\ t \geq 0 \end{matrix} \tag{7}$$

where α = liquid thermal diffusivity
r = radial coordinate within the drop
r_s = surface radius.

The subscripts r and t represent differentiation with respect to radius and time, respectively. For small droplets the assumption of spherical geometry is usually accepted. It is recognized that hwere drag exists, droplets deform; however, in this analysis the deformation is neglected. The initial condition assumes that the droplet is at uniform temperature T_o,

$$T(r,0) = T_o \qquad 0 \leq r \leq r_s \; ; \; t=0 \tag{8}$$

The boundary conditions are

$$T(0,t) \text{ is finite} \qquad r=0 \; ; \; t \geq 0 \tag{9}$$

and

$$h(T_c - T_s) = k T_r + \frac{\dot{w}}{A_d}\left[\lambda + c_{p_v}(T_c - T_s)\right] \qquad r=r_s \; ; \; 0 \leq t \leq \infty \tag{10}$$

where h = heat transfer coefficient

T_s = drop surface temperature

k = liquid thermal conductivity

λ = heat of vaporization

c_{p_v} = specific heat of droplet vapor

$\dot{w}$ = mass evaporation rate

A_d = droplet surface area

It is assumed that the film thickness surrounding the droplet is small compared to the droplet diameter.

The mass evaporation is given by

$$\dot{w} = A_d K_g p \ln \frac{p}{p-p_v} \qquad r=r_s \tag{11}$$

where p_v = droplet vapor pressure

K_g = mass transfer coefficient.

The heat and mass transfer coefficients h and K_g, respectively, are obtained from the Nusselt correlations

$$\frac{2r_s h}{k_m} = 2 + 0.6(Pr)^{1/3}(Re)^{\frac{1}{2}} \tag{12}$$

and

$$\frac{2r_s \bar{R}_u T_m K_g}{M_v D} = 2 + 0.6(Sc)^{1/3}(Re)^{\frac{1}{2}} \tag{13}$$

where k_m = vapor-gas mixture thermal conductivity

$\bar{R}_u$ = universal gas constant

M_v = molecular weight of vapor

T_m = arithmetic mean temperature, $\frac{T_c+T_s}{2}$

Pr = Prandtl number $(c_p\mu/k)_m$

Sc = Schmidt number $(\mu/D\rho)_m$

Re = Reynolds number $2r_s(VEL)(\frac{\rho}{\mu})_m$

V_d = drop velocity

D = molecular diffusion coefficient

and

$$VEL = \left((U_a-V_d)^2 + U_r^2 + U_o^2\right)^{\frac{1}{2}} \tag{14}$$

The droplet acceleration is determined by considering the momentum transfer between the liquid drop and gases due to aerodynamic drag

$$\frac{dV_d}{dt} = \frac{3}{8} C_D \frac{\rho_m}{\rho_\ell} \frac{(\Delta V)|\Delta V|}{r_s} \tag{15}$$

and the drag coefficient

$$C_D = 27 \left(\frac{2\rho_m \Delta V r_s}{\mu_m} \right)^{-.84} \tag{16}$$

where ΔV is the difference between the axial gas velocity and drop velocity.

The droplet radius is, of course, a function of time and is related to the mass evaporation rate,

$$\dot{r}_s = \frac{\dot{w}}{\rho_\ell A_d} \tag{17}$$

where $\dot{r}_s$ = surface regression rate; time rate of change of drop radius.

The analysis of droplet evaporation in a gas stream, as formulated above, is developed into a computer program for the numerical solution of the time dependent evaporation rate, droplet radius and temperature distribution within the drop. A detailed discussion of the calculation procedure and a program listing are contained in Appendix B. Beginning with a specification of the initial conditions the droplet vaporization and surface regression rates are determined from Eqs. (11) and (17). The heat balance equation (10) at the droplet surface is then used to determine the temperature gradient at the surface. By finite difference techniques the temperature gradient at a point adjacent to the surface, and the second derivative of temperature with respect to r at the surface are determined.

Equation (7) is then used to obtain the variation of surface temperature with time. Interior point temperature calculations are performed by utilizing a finite difference scheme for the solution of Eq. (7). At the end of the time interval new values are calculated for surface radius, drop velocity, surface temperature and droplet mass by integrating ghe appropriate time derivatives over the interval. The gas pressure, temperature and velocity are evaluated at the end of every interval from Eqs. (1)-(6). These thermodynamic and geometric properties then become the initial conditions for the next time interval. The procedure continues until the droplet mass is reduced to 10% of its initial value.

A summation of single-drop histories is used to determine the time variations in vaporization rate of a one-dimensional array of repetitively injected drops. An arbitrary number of drops are injected into the chamber every cycle at times distributed evenly over the oscillation period. For the case of four drops injected per cycle these would appear in the chamber at intervals of one-quarter period. Vaporization histories vary among drops injected at different times during one pressure oscillation; however, drops injected at times one period apart experience identical acoustic pressure and velocity fields and thus, have identical histories. Eventually the same number of drops are completely vaporized per cycle as are injected, and the number of drops in the array becomes constant over each full cycle. When this steady state condition is reached the fully developed array consists of a number of drops equal to the ratio of drop burning time to oscillation period times the number of drops injected per cycle. The droplets in the array have a decreasing mass down the length of the chamber and range in age from a new drop just injected to an old one almost completely vaporized. With the array fully developed, the mass vaporizing from the entire array of drops at any time is obtained from a summation of the vaporization of the individual drops that constitute the array.

Since the history of each injected drop is calculated independently of the preceding or succeeding drops, the computation time is dependent on the number of drops injected per cycle. In order to simulate a continuous injection of drops, it would be necessary to analyze an unwieldy number of drops. Alternatively, it is possible to smooth out perturbations in the array vaporization history caused by the appearance and disappearance of a small finite number of drops by artificially inserting additional drops between those whose histories are calculated. The vaporization rates of the artificially injected drops are determined by interpolation between the vaporization rates of the drops for which calculations are performed. Thus the instantaneous vaporization rate for the entire array is obtained from a summation of the calculated drop histories and interpolated artificial drop histories.

The vaporization history of eight drops, injected at equal intervals during a pressure cycle is shown in Figure 1. The vaporization rate tends to be higher at both the maximum and minimum pressure condition in the oscillation than at the mean pressure conditions. This is a result of lower total velocities at the mean pressure condition. For the case shown in Figure 1, the next drop injected (drop 9) would exhibit the identical behavior as drop 1. For a burning time of .001 seconds and a pressure oscillation period of .00033 seconds the fully developed array would consist of 30 drops with drops 1, 9, 17, 25 exhibiting identical behavior, drops 2, 10, 18, 26 exhibiting identical behavior, etc. The vaporization from the entire array is obtained by adding the evaporation rates of all of the constituent drops at corresponding times in the pressure cycle. The results of this calculation, along with the pressure curve, is shown in Figure 2. It is evident from this curve that the evaporation rate tends to be higher at times in the pressure cycle corresponding to maximum or minimum pressure with relatively lower evaporation rates occurring at the mean pressure.

The calculation shown in Figures 1 and 2 required 160 seconds of computing time on a CDC 6600 computer.

In order to insure that certain computational procedures being used would yield results that were reproducible, the following tests were conducted: In the drop evaporation test runs, calculation was terminated after 90% of the drop mass had evaporated. In order to insure that this was not a premature cutoff point, one run was extended to the 97% mass evaporation point. In the former case, the response function was 0.548; in the latter, 0.545. The difference is negligible; thus a mass termination point of 90% was adopted.

In the determination of the response factor, $R_{n\ell}$, eight drops were introduced, evenly spaced during the pressure cycle. The cumulative mass evaporation from these drops was calculated and subsequently the response factor. At low frequencies the drop array size was not statistically meaningful. The question posed was: are eight oxygen drops a statistical meaningful array at a frequency of 1000 Hertz? A quick answer was sought. In one run, 20 drops were used in a drop array and the resulting response factor thus determined did not vary significantly from the results for the eight-drop array.

RESPONSE

Imposing acoustic oscillations on the pressure and velocity field in a rocket combustion chamber causes perturbations in the vaporization rate of the array of droplets resident in the chamber. The oscillation in vaporization rate will exhibit harmonic components of the imposed frequency of acoustic oscillations. A series of response factors are calculated in this study in order to relate vaporization rate oscillations to pressure and velocity field oscillations. These response factors are the in-phase and out-of-phase components of the vaporization rate oscillations relative to the acoustic pressure oscillations, where both oscillations are given as fractional perturbations about the mean value of the variable.

The nonlinear in-phase response factor, $R_{n\ell}$, can be extracted from the array vaporization rate and normalized by correlation procedure (Ref. 5) defined by

$$R_{n\ell} = \frac{\int_0^{2\pi} W'P'd\omega t}{\int_0^{2\pi} (P')^2 d\omega t} \tag{18}$$

where

$$P' = \frac{P-\bar{P}_c}{\bar{P}_c}$$

and

$$W' = \frac{W-\bar{W}}{\bar{W}}$$

where W is the instantaneous evaporation rate from the entire array and $\bar{W}$ is the average value of W taken over a full cycle.

The nonlinear out-of-phase response factor, $I_{n\ell}$, is given by

$$I_{n\ell} = \frac{\int_0^{2\pi} W' \sum_{n=1}^{\infty} P_n \sin(2\pi f_n t - \varphi_n) d\omega t}{\int_0^{2\pi} (P')^2 d\omega t} \tag{19}$$

where

$$P_n = .859 J_1\left(\frac{1.841R}{R_w}\right) \frac{\Delta P_n}{\bar{P}_c} \tag{20}$$

Additional response factors were calculated as part of this study:

$$R_1 = \frac{\int_0^{2\pi} W' P_1 \cos(2\pi ft-\theta)\, d\omega t}{\int_0^{2\pi} (P_1 \cos(2\pi ft-\theta))^2 d\omega t} \tag{21}$$

$$R_2 = \frac{\int_0^{2\pi} W' P_2 (\cos 4\pi ft-\varphi)\, d\omega t}{\int_0^{2\pi} \left(P_2 \cos(4\pi ft-\varphi)\right)^2 d\omega t} \tag{22}$$

Out-of-phase harmonic response factors

$$I_1 = \frac{\int_0^{2\pi} W' P_1 \sin(2\pi ft-\theta)\, d\omega t}{\int_0^{2\pi} \left(P_2 \sin(2\pi ft-\theta)\right)^2 d\omega t} \tag{23}$$

$$I_2 = \frac{\int_0^{2\pi} W' P_2 \sin(4\pi ft-\theta)\, d\omega t}{\int_0^{2\pi} \left(P_2 \sin(4\pi ft-\theta)\right)^2 d\omega t} \tag{24}$$

A Princeton type "n" and "τ" defined in Ref. 6 are also calculated from the nonlinear response factors

$$\text{"n"} = \frac{R_{n\ell}(P_1^2 + P_2^2)}{P_1^2(1-\cos 2\pi f\tau)+P_2^2(1-\cos 4\pi f\tau)} \tag{25}$$

where τ is defined from the following equation:

$$\frac{P_1^2(1-\cos 2\pi f\tau)+P_2^2(1-\cos 4\pi f\tau)}{P_1^2 \sin 2\pi f\tau + P_2^2 \sin 4\pi f\tau} = -\frac{R_{n\ell}}{I_{n\ell}} \tag{26}$$

RESULTS AND DISCUSSION

A. Parametric Study of Response Functions

A series of calculations were made, utilizing the previously described computer program, in order to determine the effect of a variety of boundary conditions on the magnitude of the response functions for evaporating liquid oxygen droplets. The thermodynamic properties used in the calculations were:

Thermal diffusivity = 7.56×10^{-7} ft^2/sec
Liquid density = 72 lbm/ft^3
Prandtl number = .712
Gas viscosity = 4.2×10^{-5} lbm/ft-sec
Vapor specific heat = .288 $BTU/lbm^{\circ}F$
Liquid thermal conductivity = 2.21×10^{-5} $BTU/ft\text{-}sec\text{-}^{\circ}F$
Gas thermal conductivity = 4.04×10^{-5} $BTU/ft\text{-}sec\text{-}^{\circ}F$
Isentropic exponent = 1.135
Initial drop temperature = $140^{\circ}R$
Combustion gas temperature = $6500^{\circ}R$

The vapor pressure and heat of vaporization of the liquid oxygen is evaluated as a function of the drop surface temperature by

$$p_v(lbf/ft^2) = e^{16.93 - \frac{1476}{T-3.57}}$$

$$\lambda(BTU/lbm) = 61.33 + .5916T - .00248T^2$$

The following parameters were varied over the range indicated:

Chamber pressure: 100-600 psia
Initial drop radius: 15-150 microns
Final gas velocity: 400-2400 ft/sec
Initial drop velocity: 50-200 ft/sec
Frequency of oscillation: 200-30,000 cps
Amplitude of fundamental (Δp_1): $.01p_c$-$.8p_c$
Amplitude of harmonic (Δp_2): 0-1.2 Δp_1

Table I gives the full range of tests conducted with boundary conditions and response functions.

A cursory look at the values calculated for the in-phase nonlinear response factor, $R_{n\ell}$, indicates many values that are greater than those previously found by the methods of Refs. 2 and 4. The present study includes the effects of finite thermal conductivity which, as discussed previously, causes a significant decrease in the time required for the drop surface to reach an effective equilibrium temperature. In addition, the temperature of the drop center remains relatively unchanged over the entire lifetime of the drop with the temperature gradient in the drop concentrated at the drop outer surface. Both these factors make the drop more sensitive to fluctuations in ambient pressure and temperature with concomitant increased response factors.

The major cause of larger response factors obtained in this study, however, appears to be due primarily to the effect of wave distortion, i.e., inclusion of harmonics in the ambient pressure field. This causes more relative peaks and valleys in the impressed pressure oscillation with attendant increases in evaporation rate. Another significant contributing effect is the presence of the array of drops in a varying gas velocity field.

Figures 3 and 4 indicate the quantitative effect of wave distortion and amplitude on the nonlinear in-phase response factor, $R_{n\ell}$. For the case of a constant ratio of the first harmonic to fundamental wave pressure amplitude, $\Delta P_2/\Delta P_1=0.8$, Figure 3, the variation of $R_{n\ell}$ is relatively independent of fundamental amplitude provided $\Delta P_1 \geq 0.1P_c$. For the case where the fundamental amplitude is small, $\Delta p_1=0.01p_c$, the response factor increases significantly to a maximum value of $R_{n\ell}=1.68$. This behavior agrees qualitatively with that shown in Figure 2 of NASA TN D-6287, Ref. 5. Figures 4a, 4b and 4c show the effect of variations in the relative magnitude of the harmonic amplitude. The maximum response curve occurs at a value of $\Delta p_2=0.8\Delta p_1$. This too is in agreement with the results found in Ref. 5. It is

observed from Figures 4a,b,c that the frequency at which the maximum value of $R_{n\ell}$ occurs decreases with increasing fundamental amplitude.

Figure 4b contains the results of calculations made for zero harmonic content, $\Delta p_2=0$. As discussed above, the results obtained in Ref. 4, NASA TN D-3749, also were for the case of no harmonic content in the pressure oscillation. The magnitude of the response factors calculated in this study for $\Delta p_2=0$ are lower than those obtained with non-zero harmonics and are consistent with those obtained in NASA TN D-3749.

The final gas velocity greatly affects the value of $R_{n\ell}$. This effect is observed in Figure 5 at all frequencies. A crossplot was made at 1200 Hz, Figure 6, and the final gas velocity extended on the low scale to 50 feet per second with drop velocities of 100 feet per second. It is seen that the value of the nonlinear in-phase response factor increases to 2.36 at a final gas velocity of 200 feet per second. It is also observed that the values of $R_{n\ell}$ are about twice those observed in the previous Figures 3 and 4. In the regime $\Delta p_1 \geq 0.1 p_c$, the values and behavior of the nonlinear response factor, $R_{n\ell}$, agree both qualitatively and quantitatively with those shown in Figure 7, NASA TN D-6287, Ref. 5.

An attempt was made to correlate all of the response factor data obtained utilizing the transformed frequency suggested in Refs. 2 and 4. However, the buckshot nature of the resulting curves indicated that while qualitative agreement between the two studies is obtained, the previous correlations are unsuitable for the kind of model employed in this study. In order to correlate the data it was necessary to use a transformed response factor together with a modified transformed frequency. The transformations were obtained by trial and error in an attempt to minimize the data scatter. It was found that rather than include the effects of wave amplitude and distortion in the transformed coordinates, the effect of these variables could best be seen by utilizing parametric curves.

The transformed frequency is defined as

$$F = f\left(\frac{1200}{u_f}\right)^{1/3}\left(\frac{300}{p_c}\right)^{1/3}\left(\frac{r}{50}\right)^{3/2}$$

and the transformed response factor is

$$R_{n\ell}\left(\frac{u_f}{1200}\right)^{.56}\left(\frac{100}{V_D}\right)^{.15}\left(\frac{300}{p_c}\right)^{.1}\left(\frac{50}{r}\right)^{.12}$$

Figure 7 shows the results of the correlation study with separate curves to depict the effect of wave distortion and amplitude. For fundamental amplitudes greater than $.1p_c$ the effect of fundamental amplitude is not significant, but the effect of harmonic amplitude must be considered. Decreasing the fundamental amplitude to $0.01p_c$ causes significant increases in the response factor.

The study was extended to include the correlation of the in-phase fundamental response factor R_1 and harmonic response factor R_2, and these are shown in Figures 8 and 9. Both response factors are transformed similar to $R_{n\ell}$ and correlated against the transformed frequency factor cited above. The fundamental response factor increases with the amplitude of the harmonic and is relatively insensitive to variations in the amplitude of the fundamental for $\Delta p_1 \geq .1p_c$. The magnitude of the harmonic response function increases with a decrease in the magnitude of the harmonic pressure oscillation component. Response factors, R_2, on the order of 3.0 were calculated for harmonic amplitudes equal to 0.2 times the fundamental amplitude. The qualitative behavior, and the juxtaposition of these curves, agree with those shown in Figure 8, Ref. 5. The curves shown in Figures 7,8,9 represent the best fit through the available data. Deviations from the curve were generally less than 10% from the curve.

Figures 10,11 and 12 represent the correlation of the results for the nonlinear out-of-phase response factor, $I_{n\ell}$, the fundamental

out-of-phase response factore, I_1, and the harmonic out-of-phase response factor, I_2. The $I_{n\ell}$ curves were correlated with transformations similar to those employed for the in-phase response factor. The shape of the curve is similar to a cubic with a relative maximum point in the transformed frequency range of 300-400 cps and a relative minimum at a frequency corresponding to the location of the maximum point on the $R_{n\ell}$ curves.

The results for the correlation of the out-of-phase fundamental response factor indicate that wave amplitude and distortion have no effect. In addition, excellent correlation was obtained by transforming I_1 by a multiple of $(300/p_c)^{.1}$ and the frequency by a multiple of $(r/50)^{3/2}$. Thus the final gas velocity and initial drop velocity are not effective in varying the magnitude of I_1.

The correlation of the data for the out-of-phase harmonic response factor, I_2, required utilizing a transformed ordinate

$$I_2\left(\frac{u_f}{1200}\right)^{.2}\left(\frac{100}{V_D}\right)^{.15}\left(\frac{300}{p_c}\right)^{.1}$$

plotted against a transformed frequency

$$f\left(\frac{r}{50}\right)^{3/2}$$

Correlation also required the use of parametric curves to display the effect of wave distortion; i.e., the magnitude of I_2 decreases with increasing harmonic component amplitude.

In Figure 13, a corelation of a Princeton type "n" was plotted vs. frequency factor, F. It is seen to remain essentially constant over a broad spectrum of frequency factor. The value of "n" increases with the amplitude of the harmonic pressure perturbation amplitude.

Figure 14 is a plot of τ vs. frequency of oscillation for a variety of drop radii, initial drop velocity, final gas velocity and amplitude of pressure disturbance. The variation of the values of

of the parameters cited above are those which normally occur in liquid rocket engine practice. The results indicate that it is possible to correlate the values of τ with the frequency of oscillation without any significant dependence on any of the other parameters. The correlation seems to indicate a relationship between τ and f of the form,

$$f\tau = \text{constant}.$$

The value of τ is determined from Eq. (26). The right-hand side of the equation, i.e., $R_{n\ell}/I_{n\ell}$, has a value in the range 0.1 to 4.0 for the vast amjority of cases tested in this study. The corresponding values for $2\pi f\tau$ are in the range 250 to 350 degrees. These, when plotted for the specific case tested, yielded essentially a straight line on a log-log plot as is suggested in the above equation.

The data from the above studies were regrouped for each of the seven response factors to show the effects of fundamental pressure perturbation amplitude (Δp_1), harmonic pressure perturbation amplitude (Δp_2), injection velocity (V_D), final gas velocity (u_f), droplet radius (r) and chamber pressure (P_c). The following table shows a summary of the relative effects of the various parameters on the different response factors:

	Δp_1	Δp_2	V_D	u_f	R	P_c
$R_{n\ell}$	weak	moderate	weak	strong	strong	weak
R_1	weak	strong	weak	strong	strong	moderate
R_2	weak	strong	weak	strong	strong	weak
$I_{n\ell}$	weak	moderate	weak	moderate	moderate	weak
I_1	weak	weak	weak	weak	moderate	weak
I_2	weak	moderate	moderate	moderate	moderate	weak
n	weak	strong	moderate	strong	moderate	weak
τ	independent of all parameters					

B. Effect of Thermal Conductivity

In general, it was found that the nonlinear in-phase response factor, $R_{n\ell}$, was not very sensitive to variations of up to 10^3 in the liquid thermal conductivity, for the frequency range of interest in combustion instability studies. These results are explained below.

The calculation for the $R_{n\ell}$ requires the summation of the instantaneous evaporation rate of the complete array of drops in the chamber over an interval of time. Thus, any particular local event resulting from the interaction of the pressure wave and one drop is masked by the summation of all the other events occurring at the same time. In other words, the random event associated with each drop is masked by the stoichastic behavior of the drop array.

A second reason for the insensitivity of $R_{n\ell}$ to the thermal conductivity variation is that an open loop analysis is assumed. There is no feedback to the wave behavior from the droplet evaporation. This feedback can be significant in deforming the wave shape, and consequently the response factor, $R_{n\ell}$. The wave shape can be deformed not only by coupling the relaxation time to the evaporation but also by the amount of vaporization occurring from the array of drops during the coupling. These two situations are discussed below for a particular case.

The drop is introduced at a uniform temperature of $140^{\circ}R$. The drop surface heats up to about $240^{\circ}R$ and remains at that temperature during its lifetime. For a 50μ radius oxygen drop, and normal thermal conductivity, it takes 1.99×10^{-6} sec. for the drop surface to reach $238^{\circ}R$. For the drop with high thermal conductivity, i.e., 10^{-2} Btu $ft^{-1}{}^{\circ}R^{-1}sec^{-1}$, it takes 1.87×10^{-4} sec. for the drop surface to reach $239^{\circ}R$. It is noticed that the time lag differs by about two orders of magnitude. In a closed loop analysis, this difference in time lag could mean the difference in the effective coupling of the drop evaporation to drive the passing pressure wave.

For the case where the normal thermal conductivity is employed, the drop center does not appreciably heat up during its lifetime. For example, after 90% of the drop mass has evaporated, the temperature of the drop center is still 144^{o}R. As found previously (Ref. 3), the temperature gradient is concentrated at the drop outer surface. For the case where the large thermal conductivity is used, it is found that the drop remains uniform in temperature. The uniform temperature drop has a much larger thermal inertia than the normal drop; thus the dynamic behavior of these drops would be very different, not only in the lag times, but also in the mass evaporation response to a disturbance. For example, for a passing compression wave, the normal drop would obtain a much greater surface temperature than the uniform temperature drop for the same amount of heat transfer to the droplet. If one includes the nonlinearity of the Clausius-Clapeyron equation, then the additional mass evaporation in the former case would be much greater, with increased concomitant response factor.

SUMMARY OF RESULTS

The objective of this investigation was to analytically determine stability parameters relating drop vaporization response rates in a liquid rocket combustor to nonlinear-high amplitude pressure oscillations for drops vaporizing with finite liquid thermal conductivity. The results of the program can be summarized as follows:

1. A computer program was developed for determining the vaporization response of droplets with finite thermal conductivity to high amplitude distorted pressure oscillations.

2. Drop vaporization responses for an array of drops traveling through a rocket combustor are significantly different than the response of a single drop stationary in a flow field.

3. The vaporization rate responses were very dependent on drop size, final gas velocity in the combustor and wave distortion. The responses were moderately dependent on liquid droplet velocity and weakly dependent on chamber pressure and wave amplitude.

4. A general correlation scheme was developed to relate various response numbers that can be used in stability analyses to combustor operating conditions.

5. For the Princeton type time lag response, the time lag was found to be inversely proportional to the frequency, resulting in a phase lag of 250 to 350 degrees.

REFERENCES

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2. Heidmann, M.F. and Wieber, P.R.: "Analysis of n-Heptane Vaporization in Unstable Combustors with Travelling Transverse Oscillations" NASA TN D-3424, 1966.

3. Agosta, V.D. and Hammer, S.S.: "Droplet Vaporization with Liquid Heat Conduction". Propulsion Sciences, Inc., NASA CR-72672, 1970.

4. Heidmann, M.F. and Wieber, P.R.: "Analysis of Frequency Response Characteristics of Propellant Vaporization". NASA TN D-3749, 1966.

5. Heidmann, M.F.: "Amplification by Wave Distortion of the Dynamic Response of Vaporization Limited Combustion". NASA TN D-6287, 1971.

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7. Heidmann, M.F.: "Frequency Response of a Vaporization Process to Distorted Acoustic Disturbances". NASA TN D-6806, 1972.

TABLE I

*PARAMETRIC STUDY OF RESPONSE FUNCTIONS

*Unless otherwise noted, the boundary conditions used to calculate the response functions are:

Chamber pressure p_c = 300 psia
Drop radius = 50 microns
Drop initial temperature = 140°R
Drop initial velocity = 100 ft/sec
Final gas velocity = 1200 ft/sec
Amplitude of fundamental pressure perturbation; $\Delta p_1 = .2\ p_c$
Amplitude of harmonic pressure perturbation; $\Delta p_2 = .8\ \Delta p_1$

f	$R_{n\ell}$	R_1	R_2	$I_{n\ell}$	I_1	I_2	n	$\tau \times 10^3$
150	.289	.244	.359	.345	.252	.491	.404	5.7
200	.351	.276	.468	.442	.389	.524	.519	4.3
400	.581	.526	.666	.448	.606	.20	.541	2.019
800	.810	.910	.654	.185	.265	.0603	.538	.873
1500	.986	1.112	.789	.232	.216	.025	.65	.46
3000	.941	1.04	.781	.255	.193	.352	.682	.24
6000	.831	.987	.628	.44	.355	.583	.63	.125
12000	.549	.8	.255	.684	.609	.802	.73	.071
30000	.086	.340	-.309	.765	.808	.967	.4	.032
$\Delta p_1 = .1\ p_c$								
600	.674	.720	.602	.425	.558	.216	.555	1.3
1200	.911	.854	1.0	.131	.189	.0415	.606	.56
2400	1.026	1.032	.823	.241	.162	.411	.632	.316
5000	.844	.979	.635	.412	.329	.542	.627	.152
10000	.623	.843	.275	.630	.547	.760	.729	.084

f	$R_{n\ell}$	R_1	R_2	$I_{n\ell}$	I_1	I_2	n	$\tau x10^3$
$\Delta P_1=.4\ P_c$								
250	.431	.316	.612	.438	.466	.394	.525	3.35
500	.724	.783	.631	.331	.503	.061	.527	1.507
1200	.891	.998	.725	.160	.214	.071	.590	.57
3000	.920	1.012	.757	.240	.182	.338	.615	.236
8000	.742	.916	.470	.518	.423	.666	.647	.099
$\Delta P_1=.8\ P_c$								
400	.683	.672	.7	.322	.485	.067	.501	1.89
800	.948	1.091	.724	-.001	.003	-.0072	.674	.84
1600	.941	.979	.882	.115	.0625	.197	.628	.41
2500	.918	.995	.797	.196	.152	.265	.609	.277
6000	.814	.938	.621	.387	.299	.526	.599	.126
$\Delta P_2=0\quad \Delta P_1=.2\ P_c$								
800	.566	.566	-	.075	.075	-	.3	.67
1500	.692	.692	-	.096	.096	-	.305	.331
3000	.488	.488	-	.176	.176	-	.273	.202
6000	.43	.43	-	.32	.32	-	.335	.117
12000	.244	.244	-	.54	.54	-	.673	.072
$\Delta P_2=.2\ \Delta P_1$								
800	.791	.656	3.16	.102	.137	.322	.418	.68
1600	.874	.722	3.61	.185	.176	.399	.470	.36
3000	.72	.635	3.16	.188	.209	.580	.400	.196
6000	.655	.566	2.95	.346	.315	.792	.430	.111
12000	.455	.432	2.5	.530	.525	1.11	.550	.067

f	$R_{n\ell}$	R_1	R_2	$I_{n\ell}$	I_1	I_2	n	$\tau x10^3$
$\Delta P_2=1.2\ \Delta P_1$								
400	.548	.618	.499	.334	.716	.068	.427	2.03
800	.689	1.04	.445	.130	.295	.015	.500	.448
1500	.87	1.30	.574	.250	.193	.264	.557	.27
3000	.831	1.33	.587	.362	.302	.403	.538	.254
6000	.698	1.23	.408	.461	.371	.509	.583	.133
12000	.380	.984	.098	.710	.627	.755	.941	.076
$\Delta P_2=.2\ \Delta P_1$	$\Delta P_1=.1\ P_c$							
600	.642	.551	2.91	.411	.352	.191	.470	1.155
1500	.873	.743	4.135	.280	.291	-.005	.500	.404
4000	.821	.725	3.195	.151	.156	.472	.441	.140
10000	.519	.441	2.475	.536	.516	.665	.557	.077
$\Delta P_2=1.2\ \Delta P_1$	$\Delta P_1=.1\ P_c$							
600	.580	.788	.435	.302	.630	.075	.422	1.327
1200	.773	1.067	.568	.205	.300	.139	.644	.325
2500	.956	1.532	.62	.33	.060	.437	.719	.317
5000	.749	1.23	.410	.388	.131	.494	.544	.159
10000	.475	1.051	-.091	.664	.696	.732	.780	.088
$\Delta P_2=1.2\ \Delta P_1$	$\Delta P_1=.8\ P_c$							
400	.631	.774	.531	.034	.538	.029	.418	1.93
1000	.922	1.146	.513	.110	-.2	.128	.486	.557
2500	.831	1.098	.575	.174	-.214	.216	.487	.255
6000	.720	1.12	.539	.415	.167	.496	.512	.134
$\Delta P_2=.2\ \Delta P_1$	$\Delta P_1=.8\ P_c$							
400	.558	.498	2.06	.385	.378	.557	.428	1.76
800	.798	.722	2.69	-.041	.0752	.229	.420	.603
1600	.774	.688	2.99	+.032	-.0429	.100	.368	.322

f	$R_{n\ell}$	R_1	R_2	$I_{n\ell}$	I_1	I_2	n	$\tau x10^3$
2500	.784	.671	2.52	.098	-.0166	.119	.379	.217
6000	.688	.61	2.27	.2926	.0934	.632	.360	.106
ΔP_2=.8 ΔP_1 ΔP_1=.2 P_c U_g=400								
300	1.43	1.636	1.112	.848	1.130	.403	1.147	2.6
600	1.950	2.22	1.510	.3177	.3677	.239	1.290	1.13
1000	2.082	2.50	1.429	.090	.344	.463	1.380	.68
2000	1.720	2.12	1.340	.182	.0805	.366	1.166	.329
4000	1.645	1.941	1.181	.353	.262	.496	1.095	.1755
9000	1.390	1.755	.821	.616	.502	.786	1.007	.0834
U_g=2400								
400	.414	.322	.557	.416	.466	.326	.476	2.09
800	.608	.636	.563	.260	.391	.056	.435	.935
2000	.699	.743	.630	.086	.1	.062	.466	.332
5000	.693	.776	.562	.383	.330	.466	.537	.152
12000	.425	.603	.147	.595	.518	.715	.719	.076
U_g=400 V_D=50								
300	1.11	1.185	.994	.715	.977	.304	.926	2.62
600	1.533	1.750	1.196	.162	.262	.0056	1.028	1.098
1500	1.724	1.933	1.397	.147	.126	.180	1.164	.434
4000	1.551	1.8	1.164	.231	.178	.315	1.029	.168
9000	1.298	1.63	.766	.587	.487	.743	.941	.0836
U_g=2400 V_D=50								
400	.398	.313	.531	.430	.440	.347	.465	2.09
800	.599	.623	.561	.244	.390	.0179	.424	.929
2000	.679	.663	.575	.035	.066	-.0133	.431	.319
5000	.652	.724	.539	.359	.318	.422	.505	.154
12000	.406	.578	.138	.591	.518	.703	.724	.073

f	$R_{n\ell}$	R_1	R_2	$I_{n\ell}$	I_1	I_2	n	$\tau x 10^3$
U_g=400 V_D=200								
600	2.415	2.876	1.694	.254	.246	.266	1.619	1.097
1000	2.00	2.59	1.064	.145	.075	.255	1.355	.648
2000	1.805	2.036	1.438	-.026	-.047	.0057	1.275	.191
4000	1.707	1.96	1.35	.135	.06	.253	1.155	.168
9000	1.5	1.86	.937	.515	.382	.723	1.03	.0809
U_g=2400 V_D=200								
800	.716	.868	.478	.374	.402	.332	.544	.958
2000	.779	.939	.528	.237	.101	.451	.527	.359
5000	.700	.758	.610	.339	.2811	.430	.518	.152
12000	.442	.633	.148	.601	.523	.721	.718	.072
U_g=1200 V_D=200								
200	.441	.331	.612	.499	.459	.564	.578	4.25
800	1.056	1.087	1.008	.209	.308	.0557	.7	.862
1500	1.22	1.44	.867	.1482	.519	.425	.858	.494
3000	1.019	1.2	.735	.288	.131	.535	.685	.237
6000	.935	1.101	.675	.425	.334	.567	.679	.125
12000	.586	.85	.173	.695	.621	.816	.810	.071
U_g=1200 V_D=50								
200	.321	.253	.426	.416	.354	.511	.494	4.32
800	.733	.82	.596	.161	.223	.0646	.486	.87
1500	.825	.889	.724	.084	.113	.037	.554	.44
3000	.923	1.004	.795	.323	.305	.350	.635	.243
6000	.79	.928	.574	.452	.384	.557	.622	.129
12000	.508	.738	.149	.667	.597	.777	.791	.072

f	$R_{n\ell}$	R_1	R_2	$I_{n\ell}$	I_1	I_2	n	$\tau \times 10^3$
$r=15\mu$ $U_q=1200$ $V_D=100$								
2000	.431	.340	.573	.473	.507	.428	.551	.423
4000	.660	.713	.577	.374	.409	.162	.518	.193
7000	.792	.822	.745	.197	.225	.153	.528	.1
15000	.811	.914	.627	.474	.18	.538	.641	.05
30000	.578	.781	.261	.600	.325	.719	.695	.028
$r=150\mu$								
150	.888	.990	.730	.168	.259	.0266	.589	4.58
300	1.111	1.25	.913	.299	.17	.228	.747	2.31
600	1.052	1.217	.793	.462	.297	.563	.758	1.25
1200	.822	1.029	.497	.575	.501	.691	.718	.664
3000	.261	.565	-.233	.740	.718	.665	1.342	.311
$r=50\mu$ $p_c=100$								
500	.712	.797	.580	.195	.258	.096	.477	1.42
1000	.853	.967	.676	.162	.172	.147	.565	.687
2000	.870	.953	.741	.191	.165	.232	.577	.348
4000	.783	.897	.603	.281	.211	.390	.541	.183
8000	.629	.799	.362	.523	.429	.669	.620	.102
$p_c=600$								
500	.568	.498	.677	.501	.637	.296	.587	1.65
1000	.847	.959	.672	.255	.348	.109	.572	.718
2000	1.05	1.167	.866	.3	.285	.325	.706	.357
4000	.974	1.105	.769	.343	.262	.469	.671	.183
8000	.803	1.00	.493	.567	.475	.711	.706	.099
16000	.401	.711	-.083	.792	.760	.842	1.119	.056

Symbols

a	speed of sound
A_d	droplet surface area
C_D	drag coefficient
C_{pv}	droplet vapor specific heat
D	molecular diffusion coefficient
f	frequency
h	film heat transfer coefficient
k	liquid thermal conductivity
k_m	vapor-gas mixture thermal conductivity
K_g	mass transfer coefficient
M	molecular weight
m	droplet mass
p_c	mean chamber pressure
P_v	vapor pressure
Δp_1	amplitude of fundamental pressure perturbation
Δp_2	amplitude of harmonic pressure perturbation
r	droplet radius
$\dot{r}_s$	regression rate of droplet surface
Re	Reynolds number
t	time
T_c	chamber temperature
T_s	droplet surface temperature
U	gas velocity
V_d	drop velocity
$\dot{w}$	mass evaporation rate
α	thermal diffusivity
γ	isentropic exponent
ρ	density
λ	heat of vaporization
μ	viscosity

APPENDIX B

Calculation Procedure and Program Listing

The procedure used in calculating the droplet evaporation histories, temperature distribution within the drop and response factors is illustrated by the flow diagram.

Step 1. Load into the machine the following boundary conditions, initial conditions and computational parameters.

Card 1. (14 I5)

- NRUN - Number of test case
- NMONTH - Month
- MDAY - Day
- MYEAR - Year
- JA - Number of calculation steps between output for each drop
- NA - Number of mesh points minus one within drop
- NP - Number of drops injected per cycle
- NY - Number of summation histories per period
- NART - Number of artificial drops insisted between each of tne NP calculated drops

Card 2. (8E 10.4)

- S - Initial drop radius (ft.)
- PO - Mean chamber pressure (lbf/ft^2)
- VGAF - Final gas velocity (ft/sec)
- VDI - Initial drop velocity (ft/sec)
- DPC - Ratio of peak-to-peak fundamental pressure oscillation to mean chamber pressure
- DPCI - Ratio of harmonic pressure oscillation to fundamental oscillations
- OMEGA - Frequency of pressure oscillations (cps)
- A - Stretching parameter (1.3)

Card 3. (8E 10.4)	STAB	- Initial time increment (sec.)
	TFO	- Mean combustion gas temperature (^{o}R)
	TO	- Initial droplet temperature (^{o}R)
	THETA	- Phase angle of pressure oscillation
	AJ_0	- Zero order Bessel function $J_0(\frac{1.841R}{Rw})$
	AJ_1	- First order Bessel function $J_1(\frac{1.841R}{Rw})$
	AJ_2	- Second order Bessel function $J_2(\frac{1.841R}{Rw})$
	RWR	- Ratio of radial location in chamber to wall radius
Card 4. (8E 10.4)	GAMMA	- Ratio of specific heats for combustion gases
	ALPHA	- Liquid thermal diffusivity (ft^2/sec)
	RHOL	- Liquid density (lbm/ft^3)
	RR	- Universal gas constant
	PCA	- Critical pressure - droplet (lbf/ft^2)
	PCB	- Critical pressure - combustion gases (lbf/ft^2)
	TCA	- Critical temperature - droplet (^{o}R)
	TCB	- Critical temperature - combustion gases (^{o}R)
Card 5. (8E 10.4)	EMA	- Molecular weight - droplet
	EMB	- Molecular weight - gases
	AA	- Constant in diffusivity equation - 2.33×10^{-6}
	BB	- Constant in diffusivity equation - 1.823
	EMV	- Molecular weight of drop vapor
	PR	- Prandtl number of gases
	AKB	- Combustion gas thermal conductivity ($BTU/ft\text{-}sec\text{-}^{o}R$)
	CPV	- Drop vapor specific heat
Card 6.(8E 10.4)	VIS	- Combustion gas viscosity (lbm/ft-sec)
	AK	- Drop thermal conductivity ($BTU/ft\text{-}sec\text{-}^{o}R$)

Step 2. Write out the value of all input variables and calculate all parameters that are constant throughout calculations. Initialize droplet and gas parameters to begin calculation of N^{th} drop history. Redefine variables as follows:

$$U = Tr \tag{A1}$$

so that Eq. (7) becomes

$$U_t = \alpha U_{tt} \tag{A2}$$

The space variable within the drop is also redefined:

$$a = r/s \tag{A3}$$

where s is the drop surface radius as a function of time. The space variable is further redefined as:

$$X = A\sigma\left(z+1/(A-z)\right) - \sigma \tag{A4}$$

wh where

$$\sigma = (A-1)/(A^2+1-A) \tag{A5}$$

and A is a stretching parameter which concentrates mesh points near the surface of the drop. The heat conduction equation (A2) can now be written:

$$U_T = \left(\frac{\alpha S_t z Y}{s} + \alpha Y Y_x/s^2\right)U_x + \frac{\alpha Y^2}{s^2} U_{xx} \tag{A6}$$

where $Y = \frac{dx}{dz}$.

Step 3. The interior point calculations are performed (temperature distribution within the drop) by utilizing a finite difference scheme for the solution of the transformed heat conduction equation based upon the current values of drop surface temperature and heat transfer rate to the drop.

Step 4. Droplet evaporation rates, changes in combustion gas properties, droplet velocity and net heat transfer to the drop are calculated by solving Eqs. (10) through (17). The evaporation rate is stored for future summation

in the array history. Intermediate results are printed at the end of every JA time increments.

Step 5. If the drop is not 90% evaporated, time and other thermodynamic and dynamic variables are incremented and control is transferred to Step 3.

Step 6. If the drop has reduced to 10% of its initial mass, the evaporation rate is transferred into the array summation matrix at a predetermined number of points during the period (NY).

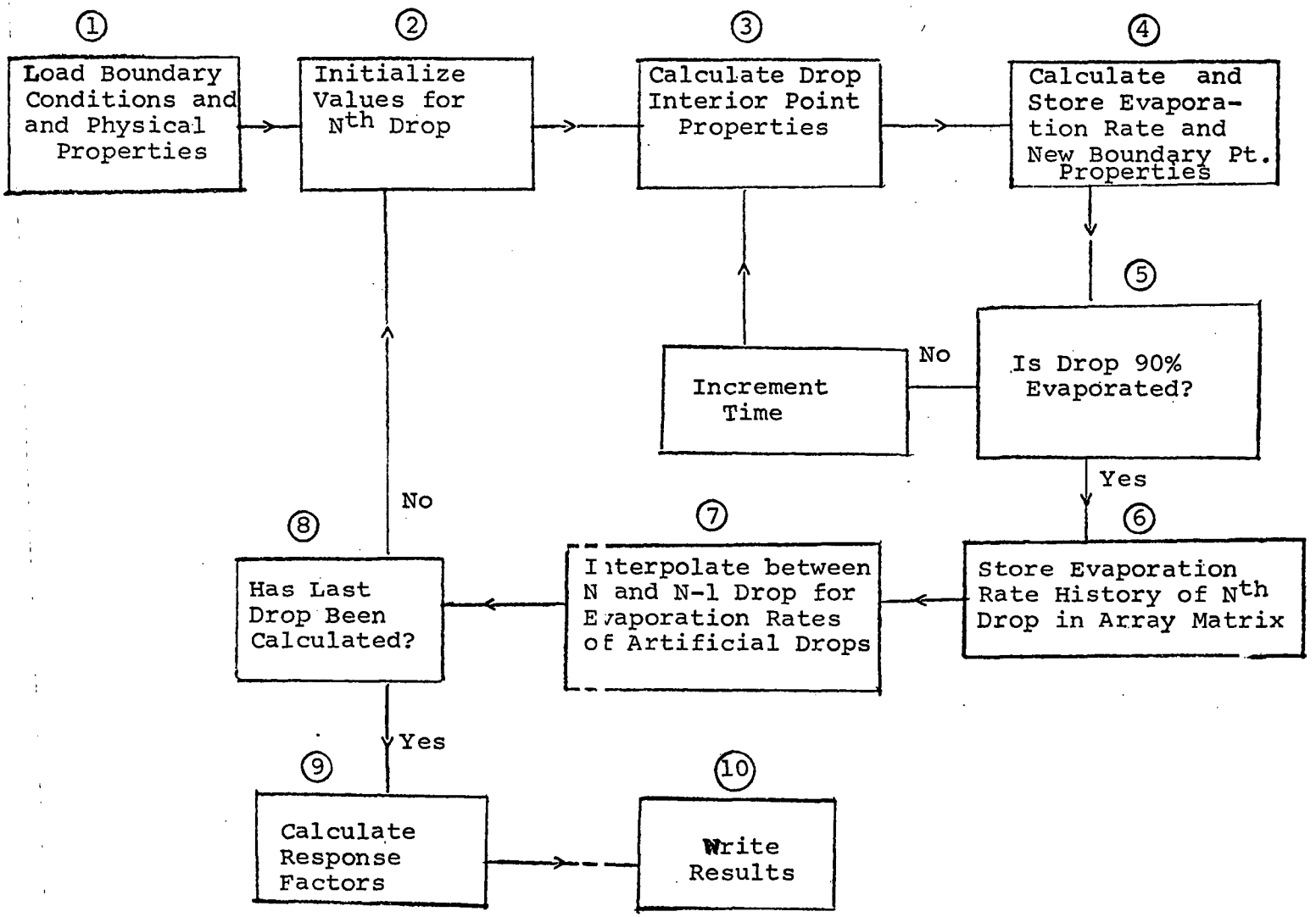
Step 7. An interpolation of the evaporate rate between the N^{th} and $(N-1)^{th}$ drop is made for each of the artificial drops injected between two drops for which calculations are performed. These evaporation rates are added to the array matrix.

Step 8. Test for end of evaporation calculations.

Step 9. Calculation of response factors by solving Eqs. (18) through (26).

Step 10. Print results.

The machine computation time is primarily a function of the number of drops injected per cycle. For the case described in Figs. 1 and 2, i.e., eight drops injected per cycle, a total of 160 seconds of CDC 6600 computer time was required.


1
Load Boundary Conditions and and Physical Properties
2
Initialize Values for Nth Drop
3
Calculate Drop Interior Point Properties
4
Calculate and Store Evaporation Rate and New Boundary Pt. Properties
5
Is Drop 90% Evaporated?
No
Increment Time
Yes
6
Store Evaporation Rate History of Nth Drop in Array Matrix
7
Interpolate between N and N-1 Drop for Evaporation Rates of Artificial Drops
8
Has Last Drop Been Calculated?
No
Yes
9
Calculate Response Factors
10
Write Results

FLOW CHART

```
                PROGRAM DROPS(INPUT,OUTPUT,TAPE5 = INPUT,TAPE6 = OUTPUT)
      C
      C         PROGRAM DROPS WITH RNL,INL,R1,R2,I1,I2,F,G, AND N AND TAU  AUG. 7
      C
000003            COMMON WWW(3000),TTT(3000),W(400),INDEX,KK2,VEL,VD,AKG,DD,PPV,H,D
     1M1,SIT,S,TIME,J,K,JA,NA,NC,DX,DT,ST,SN,STN,WDOT,P,TF
000003            COMMON SAVW(30,400),MARK,KK,NART,NN,NP
000003            COMMON X(64),Z(64),AB(64),AC(64),AD(64),U0(64),U(64),UN(64),T(64)
     1R(64)
      C
000003            PV(W) = 2980.9579867*EXP(8.928-1476.5/(W-3.568))
000016            SL(W) = 61.33+.5916*W-.00248*W**2
000025            D(W,O) = DD1/W*AA*(O/DD2)**BB
      C
      C         NA=NUMBER OF MESH POINTS MINUS ONE
      C         JA=NUMBER OF STEPS BETWEEN OUTPUTS FOR THE FIRST DROP
      C         NP=NUMBER OF DROPS INJECTED PER CYCLE
      C         NY=NUMBER OF SUMMATION HISTORIES PER PERIOD (400 MAXIMUM)
      C         A=STRETCHING PARAMETER
      C
000036            READ(5,250) NRUN,MONTH,MDAY,MYEAR,JA,NA,NP,NY,NART
000064            READ(5,260) S,P0,VGAF,VDI,DPC,DPC1,OMEGA,A
000110            READ(5,260) STAB,TF0,T0,THETA,AJ0,AJ1,AJ2,RWR
000134            READ (5,260) GAMMA,ALPHA,RHOL,RR,PCA,PCB,TCA,TCB
000160            READ (5,260) EMA,EMB,AA,BB,EMV,PR,AKB,CPV
000204            READ(5,260) VIS,AK
000214            DD2 = SQRT(TCA*TCB)
000221            DD1 = (PCA*PCB)**.333333/SQRT(EMA*EMB/(EMA+EMB))*(TCA*TCB)**.4166
     17
000241            TAV = (TF0+T0)*.5
000244            ROAV = P0*EMB/(RR*TAV)
000247            SCH = VIS/ROAV/D(P0,TAV)
000253            SIT = S
000254            NC = NA+1
000256            TAU = 1./OMEGA
000260            NX = NY-1
000262            DTST = TAU/FLOAT(NX)
000264            DTBAR = TAU/FLOAT(NP)
000266            WRITE (6,320)
000271            WRITE(6,330)NP,NART,NY
000303            WRITE(6,270)NRUN,MONTH,MDAY,MYEAR,JA,P0,TF0,S,T0,VGAF,VDI
000335            WRITE(6,340) OMEGA,THETA,DPC,AJ0,AJ1,AJ2,RWR
000357            WRITE (6,280) ALPHA,RHOL,RR,PCA,PCB,TCA,TCB,EMA,EMB,EMV,AA,BB,SCH
     1PR,AKB,CPV,AK,GAMMA,VIS
000431            WRITE (6,290)
000435            WRITE (6,300) NA,A,STAB
000447            WRITE(6,410)DPC1
      C
      C         CONVERT DPC TO 1/2 PEAK TO PEAK PERTURBATION
      C
      C         CONVERT DPC1 TO RATIO OF DP1 TO P0
      C
000455            DPC=DPC/2.
000457            DPC1=DPC*DPC1
000460            PI = 4.*ATAN(1.)
000464            DUM8=PI*4.*RHOL
000466            PI2 = 2.*PI
```

```
000467          PI02 = PI/2.
000471          POMEGA = PI2*OMEGA
000473          GA = (GAMMA-1.)/GAMMA
000475          VD = VDI
000477          TI = 0.
000500          CC1=DPC*(AJ0-AJ2)*.430/GAMMA
000504          CC2=RWR*AJ1*.467*DPC/GAMMA
000510          DUM4 = EMV/2./RR
000513          DX = 1./FLOAT(NA)
000515          DDX = 1./DX
000516          DDX2 = 1./DX/DX
000520          TWODX=2.*DX
000521          SIG = (A-1.)/(A*(A-1.)+1.)
000526          A1 = A*SIG
000527          A2 = 2.*A1
000530          A3 = SIG*(A**2+1.)
000533          A4 = A2**2/SIG
000534          DO 1 I=1,30
000536          DO 1 J=1,400
000537        1 SAVW(I,J)=0.
          C
          C     INITIALIZATION OF VALUES FOR THE (NN)TH DROP
          C
000547          DO 220 NN = 1,NP
000550          S = SIT
000552          S2 = S**2
000553          SN = S
000554          WDOT = 0.
000555          STS = 0.
000556          ST = 0.
000557          K = 0
000560          J = 0
000561          DT = STAB
000562          TIME = DTBAR*(NN-1)
000566          KK1 = 0
000567          KK2 = 0
000570          VD = VDI
000571          P=P0*(1.+.859*DPC*AJ1*COS(POMEGA*TIME-THETA)+.859*DPC1*AJ1*COS(2.
               1POMEGA*TIME))
000613          TF = TF0*(P/P0)**GA
000620          TAV = (TF0+T0)*.5
000623          ROAV = P0*EMB/(RR*TAV)
000626          SCH = VIS/ROAV/D(P,TAV)
000632          DUM2 = .6*PR**.333333
000636          DO 10 N = 1,NC
000637          X(N) = DX*FLOAT(N-1)
000642          Z(N) = (A3+X(N)-SQRT((A3+X(N))**2-A4*X(N)))/A2
000656          R(N) = Z(N)*S
000660          F = X(N)/A1+1./A-Z(N)
000665          F2 = F*F
000666          F3 = F*F2
000667          Y = A1*(1.+F2)
000672          AB(N) = Z(N)*Y
000675          AC(N) = A2*F3*ALPHA
000700          T(N) = T0
000701          IF (N.EQ.NA) YNA = Y
000704          AD(N) = Y**2*ALPHA
000707          U(N) = R(N)*T0
```

```
000711        10 CONTINUE
000718           TINT = TIME
                 C
                 C     TIME DEPENDENT COMPUTATION BEGINS HERE
                 C
000715        20 IF(NN.EQ.1.AND.J.EQ.JA)CALL OUTPUT
000726           K = K+1
000730           J = J+1
000731           IF(K.LT.100) GO TO 40
000733           DT=STAB*10.
000735           IF(K.LT.200) GO TO 40
000737           DT = 100.
000741           DO 30 N = 1,NA
000742           DT3 = (R(N+1)-R(N))**2/ALPHA/4.
000746           IF (DT.LT.DT3) GO TO 30
000751           DT = DT3
000751        30 CONTINUE
000754        40 TIME = TIME+DT
000756           UN(1) = 0.
                 C
                 C     INTERIOR POINT COMPUTATION BEGINS HERE
                 C
000757           LOOP = 0
000760        50 DO 70 N = 2,NA
000762           NQ = N+LOOP
000764           NM = NQ-1
000766           UX = (U(NQ)-U(NM))*DDX
000771           UXX = (U(N+1)+U(N-1)-2.*U(N))*DDX2
000776           AE = AB(N)*STS+AC(N)/S2
001002           AG = AD(N)/S2
001004           UT = AE*UX+AG*UXX
001007           IF (LOOP.EQ.1) GO TO 60
001011           UN(N) = U(N)+UT*DT
001015           GO TO 70
001016        60 UN(N) = (UO(N)+U(N)+UT*DT)*.5
001024        70 CONTINUE
                 C
                 C     BOUNDARY POINT COMPUTATION BEGINS HERE
                 C
001027           DT1 = DT
001030           TIM = 0.
001031           TI = TIME-DT
001032           KIM=30
001033           DT1 = DT/FLOAT(KIM)
001035        80 CONTINUE
001035           IF (LOOP.EQ.0) SN = S+ST*DT
001041           IF (LOOP.EQ.1) SN = .5*(S+SO+ST*DT)
001050           TNAMN = UN(NA-1)/SN/Z(NA-1)
001053           IF (LOOP.EQ.0) GO TO 90
001054           T(NC) = UO(NC)/SO
001057           U(NC) = UO(NC)
001060           S = SO
001061        90 TI=TIME-DT/2.
001064           P=P0*(1.+.859*DPC*AJ1*COS(POMEGA*TI-THETA)+.859*DPC1*AJ1*COS(2.*P(
                1MEGA*TI))
001106           TF = TF0*(P/P0)**GA
001113           TAV = (TF+T(NC))*.5
001116           DD = D(P,TAV)
```

```
001120          ROAV = P*EMB/(RR*TAV)
001123          VOR = VIS/ROAV
001125          DUM1 = .6*(VOR/DD)**.333333
001132          AS = SQRT(GAMMA*32.2*P/ROAV)
001140          VR=CC1*AS*(COS(POMEGA*TI+PID2-THETA)+DPC1/DPC*COS(2.*POMEGA*TI+PI
               12))
001161          VH=CC2*AS*(COS(POMEGA*TI-THETA)+DPC1/DPC*COS(2.*POMEGA*TI))
001200          VGA = VGAF*(1.-(S/SIT)**3)
001204          DV = VGA-VD
001206          VT = 5.65*(ROAV*ABS(DV))**.16*(VIS/S)**.84*DV/RHOL/S
001224          VD=VD+VT*DT/2.
001230          VEL=SQRT(DV**2+VR**2+VH**2)
001237          DOM1 = SQRT(2.*VEL*S/VOR)
001244          H = AKB/2./S*(2.+DUM2*DOM1)
001251          AKG = (2.+DUM1*DOM1)*DD*DUM4/S/TAV
001257          AKGP=AKG*P
001261          S1=S
001262       95 PPV=PV(T(NC))
001266          DUM=1.-PPV/P
001271          IF (DUM.GT.1.E-20) GO TO 100
001274          WRITE (6,310)
001277          CALL OUTPUT
001300          CALL EXIT
001301      100 DUM=AKGP*ALOG(DUM)
001305          STN = DUM/RHOL
001307          DUM = -DUM
001310          TR = -((T(NC)-TF)*H-(T(NC)-TF0)*DUM*CPV+DUM*SL(T(NC)))/AK
001325          TNAM = T(NA-1)+(TNAMN-T(NA-1))*TIM/DT
001332          TX=(T(NC)-TNAM)/TWODX
001336          SIM =S1+ST*DT1
001341          RNA= S1*Z(NA)
001343          TRNA = TX*YNA/S1
001345          TRR = (TR-TRNA)/(S1-RNA)
001351          UR = T(NC)+S1*TR
001355          UT = ALPHA*(S1*TRR+2.*TR)+UR*ST
001363          UN(NC) = U(NC)+UT*DT1
001367          T(NC) = UN(NC)/SIM
001371          ST = STN
001372          S1= SIM
001373          TIM = TIM+DT1
001375          IF(TIM.LT.DT)GO TO 95
001377          IF (KIM.GT.0) SN = SIM
001402          DO 110 N = 1,NC
001404          UO(N) = U(N)
001406          R(N) = SN*Z(N)
001410          IF (N.NE.1) T(N) = UN(N)/R(N)
001414      110 U(N) = UN(N)
001420          S0 = S
001422          S = SN
001423          S2 = S**2
001424          ST = STN
001425          STS = ST/S
001426          IF (LOOP.EQ.1) GO TO 120
001430          LOOP = 1
001431          GO TO 50
001432      120 CONTINUE
            C
            C       INTERIOR POINT CALCULATION AND BOUNDARY POINT CALCULATION END HERE
```

```
      C
001432        WDOT = DUM8*S2*ST
001435        KK1 = KK1+1
001436        IF (KK1.LT.3) GO TO 130
001440        KK2 = KK2+1
001441        KK1 = 0
001442        WWW(KK2) = -WDOT
001444        TTT(KK2) = TIME
001445    130 CONTINUE
001445        T(1)=(UN(2)*Z(3)**2/Z(2)-UN(3)*Z(2)**2/Z(3))/(Z(3)**2-Z(2)**2)/SN
001456        IF (NN.EQ.1) GO TO 140
001460        RATIO = (S/SIT)**3
001463        IF (RATIO .GE. 0.1 ) GO TO 20
001465        TB=TIME-TINT
001467        WRITE (6,350) NN,RATIO
001477        GO TO 150
001500    140 IF (S.GT.0.4641589*SIT) GO TO 20
001505        WRITE (6,360)
001510        TB = TIME
001512        IF (NN.EQ.1)  T90 = TB
001515    150 CONTINUE
      C
      C     COMPUTATION OF SUMMATION HISTORIES BEGINS HERE
      C
001515        DUMA1=TB+DTBAR*(NN-1)
001523        DUMA2=TAU+DTBAR*(NN-1)
001527        IF(TB.GE.TAU)GO TO 155
001531        MARK=1
001532        DO 153 KK=1,NY
001533        IF(NN.EQ.1)W(KK)=0.
001536        TK=TB+(KK-1)*DTST
001545        IF(TK.GT.DUMA1.AND.TK.LT.DUMA2)TK=-100.
001554        IF(TK.GT.TB+DUMA2)TK=-100.
001560        IF(TK.GE.DUMA2)TK=TK-TAU
001565        INDEX=KK2
001567        W(KK)=FWDOT(TK)+W(KK)
001574    153 CONTINUE
001577        GO TO 220
001577    155 DO 210 KK=1,NY
001601        MARK=1
001602        TK = TB+(KK-1)*DTST
001607        I = 1
001610    160 TI = TB+(I-1)*DTBAR
001616        IF (TK.GT.TI) GO TO 170
001621        II = I
001622        GO TO 180
001622    170 I = I+1
001624        GO TO 160
001624    180 CONTINUE
001624        IF (NN.EQ.1) W(KK) = 0.
001630        INDEX = KK2
001632        DO 190 N = 1,2
001633        IF (NN.LT.II.AND.N.EQ.1) GO TO 190
001642        JJ = N-1
001643        TI = TK-JJ*TAU
001646        W(KK) = W(KK)+FWDOT(TI)
001653        MARK=MARK+1
001654    190 CONTINUE
```

```
001656          NT = TK/TAU
001661          DO 200 NI = 2,NT
001663          T2 = NI*TAU+(NN-1)*DTBAR
001671          IF (TK.LT.T2) GO TO 200
001674          TI = TK-NI*TAU
001677          W(KK) = W(KK)+FWDOT(TI)
001704          MARK=MARK+1
001705      200 CONTINUE
001710      210 CONTINUE
001712      220 CONTINUE
001715          WRITE (6,380)
001720          WRITE (6,370) (W(KK),KK = 1,NY)
001733          WAV = 0.
001734          DO 230 KK = 1,NX
001736      230 WAV = WAV+(W(KK)+W(KK+1))/2.*DTST
001746          WAV = WAV/TAU
001747          WRITE (6,390) WAV
          C
          C             COMPUTATION OF RESPONSE FACTORS BEGINS HERE
          C
001755          PSQ=0.
001756          P1SQ=0.
001757          P2SQ=0.
001760          PISQ=0.
001761          PI1SQ=0.
001762          PI2SQ=0.
001763          RNL=0.
001764          R1=0.
001765          R2=0.
001766          AINL=0.
001767          AI1=0.
001770          AI2=0.
001771          DO 231  KK=1,NY
001772          TK=TB+(KK-1)*DTST
001777          PM=1.
002001          IF(KK.EQ.1.OR.KK.EQ.NY) PM=.5
002011          APSQ=.859*AJ1*(DPC*COS(POMEGA*TK-THETA)+DPC1*COS(2.*POMEGA*TK))
002030          AP1SQ=.859*AJ1*DPC*COS(POMEGA*TK-THETA)
002040          AP2SQ=.859*AJ1*DPC1*COS(2.*POMEGA*TK)
002050          APISQ=.859*AJ1*(DPC*SIN(POMEGA*TK-THETA)+DPC1*SIN(2.*POMEGA*TK))
002067          API1SQ=.859*AJ1*DPC*SIN(POMEGA*TK-THETA)
002077          API2SQ=.859*AJ1*DPC1*SIN(2.*POMEGA*TK)
002107          WPRM=(W(KK)-WAV)/WAV
002112          RNL=RNL+PM*WPRM*APSQ
002115          R1=R1+PM*WPRM*AP1SQ
002120          R2=R2+PM*WPRM*AP2SQ
002123          AINL=AINL+PM*WPRM*APISQ
002126          AI1=AI1+PM*WPRM*API1SQ
002131          AI2=AI2+PM*WPRM*API2SQ
002134          PSQ=PSQ+PM*APSQ**2
002137          P1SQ=P1SQ+PM*AP1SQ**2
002143          P2SQ=P2SQ+PM*AP2SQ**2
002147          PISQ=PISQ+PM*APISQ**2
002153          PI1SQ=PI1SQ+PM*API1SQ**2
002157      231 PI2SQ=PI2SQ+PM*API2SQ**2
002166          RNL=RNL/PSQ
002167          R1=R1/P1SQ
002171          R2=R2/P2SQ
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002172          AINL=AINL/PISQ
002174          AI1=AI1/PI1SQ
002175          AI2=AI2/PI2SQ
002177          CAPF=OMEGA*(SIT/.000166)**1.5*(800./VGAF)**.3333*(.1/DPC)**.3333*
               1(43200./P0)**.3333
002225          CAPG=CAPF*(VGAF/800.)**(7./3.)*(DPC/.1)**.633
002243          WRITE(6,400)RNL, R1,R2,AINL,AI1,AI2,CAPF,CAPG,T90
          C
          C          CALCULATION OF N AND TAU BEGINS HERE
          C
002270          AP=.859*AJ1*DPC
002273          BP=.859*AJ1*DPC1
002275          WT=PI2/2000.
002277          SAVFX=0.
002300          SAVJ=0.
002301          DO 240 I=1,2000
002302          FX=RNL/AINL*(AP**2*SIN(WT)+BP**2*SIN(2.*WT))+BP**2*(1.-COS(2.*WT))
               1+AP**2*(1.-COS(WT))
002335          IF(FX) 232,238,233
002336       232 J=-1
002337          GO TO 234
002340       233 J=1
002341       234 IF(J+SAVJ)239,238,239
002344       238 WTS=WT+PI2/2000.*(FX/(SAVFX-FX))
002352          PTAU=WTS/POMEGA
002354          ANGLE=WTS*360./PI2
002356          PEN=RNL*(AP**2+BP**2)/(AP**2*(1.-COS(WT))+BP**2*(1.-COS(2.*WT)))
002377          WRITE(6,500) WTS,ANGLE,PTAU,PEN
002413       239 WT=(I+1)/2000.*PI2
002420          SAVFX=FX
002421       240 SAVJ=J
002425          CALL EXIT
          C
002426       250 FORMAT (14I5)
002426       260 FORMAT (8E10.4)
002426       270 FORMAT (1H0, 10HRUN NUMBERI5,5X,2HONI3,1H/I2,1H/I2//
               1                                   13H OUTPUT EVERYI6,6H STEPS//14H GAS
               2 PRESSURE=E13.5,7H LB/FT2/17H GAS TEMPERATURE=E13.5,8H RANKINE/24H
               3 DROPLET INITIAL RADIUS=E13.5,3H FT/21H DROPLET TEMPERATURE=E13.5,
               48H RANKINE/11H FINAL VGA=E13.5,7H FT/SEC/18H DROPLET VELOCITY=E13.
               55,7H FT/SEC/51H AUTOMATIC STOP WHEN 90 PERCENT MASS HAS EVAPORATED
               6)
002426       280 FORMAT (9H0   ALPHA,7X,4HRHOL,6X,12HGAS CONSTANT,3X,3HPCA,9X,3HPCB
               1,9X,3HTCA,9X,3HTCB/7E12.4//4X,2HMA,10X,2HMB,10X,2HMV,11X,1HA,11X,1
               28,9X,3HSCH,9X,2HPR/7E12.4//4X,2HKB,8X,5H(CP)V,7X,5HKAPPA,7X,5HGAMM
               3A,7X,3HVIS/5E12.4/)
002426       290 FORMAT (35H PV(T)=EXP(16.928-1476.5/(T-3.568))/36H LAMBDA(T)=61.33
               1+.5916*T-.00248*T**2/83H D(P,T)=(PCA*PCB)**1/3/SQRT(MA*MB/(MA+MB))
               2*(TCA*TCB)**5/12/P*A*(T/SQRT(TCA*TCB))**B)
002426       300 FORMAT (36HONUMBER OF INTERVALS INSIDE DROPLET=I3,23H, STRETCHING
               1PARAMETER=E12.4/21H STABILITY PARAMETER=E12.4)
002426       310 FORMAT (6HOFAIL1)
002426       320 FORMAT(1H1, 36HPROGRAM DROPS AUGUST 1972 VERSION            )
002426       330 FORMAT(46H THE NUMBER OF CALCULATED DROPS PER PERIOD IS I5/58H THE
               1 NUMBER OF ARTIFICIAL DROPS BETWEEN EACH REAL DROP IS I5/51H THE E
               2VAPORATION FROM THE ENTIRE ARRAY IS SUMMED AT,I4,18H POINTS PER PE
               3RIOD)
002426       340 FORMAT(9H0   OMEGA,7X,5HTHETA,5X,10HDPC  PK-PK,6X,3HAJ0,9X,3HAJ1,
```

```
     19X,3HAJ2,9X,3HRWR/7E12.4)
002426     350 FORMAT(1H0,5HDROP I5,36H CALCULATION ENDS WITH A MASS RATIO=E10.3)
002426     360 FORMAT (1H0,5X,32H90 PERCENT EVAPORATION COMPLETED)
002426     370 FORMAT (1X,10E10.3)
002426     380 FORMAT (1H1,6H  W IS)
002426     390 FORMAT(1X,5H WAV= E12.5)
002426     400 FORMAT(1X,5H RNL= E12.5/1X,5H R1= E12.5/1X,5H R2= E12.5/1X,5H INL=
     1 E12.5/1X,5H I1= E12.5/1X,5H I2= E12.5/1X,5H F = E12.5/1X,5H G =
     2E12.5/1X,5H T90=E12.5, 8H SECONDS)
002426     410 FORMAT(1X,5HDPC1=E12.5,5X,9H(DP1/DPC) )
002426     500 FORMAT(/// 9H WTS =   E12.5,8H RADIANS/9H ANGLE = E12.5,8H DEGREES
     1/9H TAU =   E12.5,8H SECONDS/9H N =     E12.5)
002426         END
```

```
                FUNCTION FWDOT (T)
000003          COMMON WWW(3000),TTT(3000),W(400),INDEX,KK2,VEL,VD,AKG,DD,PPV,H,DO
               1M1,SIT,S,TIME,J,K,JA,NA,NC,DX,DT,ST,SN,STN,WDOT,P,TF
000003          COMMON SAVW(30,400),MARK,KK,NART,NN,NP
          C
000003          IF(T.LT.TTT(1))FWDOT=0.
000006          IF(T.LT.TTT(1)) GO TO 30
000010          IF (T    .GT.TTT(KK2)) FWDOT = WWW(KK2)
000015          IF(T.GT.TTT(KK2)) GO TO 30
000021          DO 10 I = 1,INDEX
000022          L = INDEX+1-I
000024          IF (TTT(L).GE.T    ) GO TO 10
000027          LM = L
000030          GO TO 20
000031       10 CONTINUE
000034       20 LP = LM+1
000036          INDEX = LP
000037          EPS = (TTT(LM)-T    )/(TTT(LM)-TTT(LP))
000044          FWDOT = WWW(LM)+EPS*(WWW(LP)-WWW(LM))
          C
          C     BEGINNING OF ARTIFICAL DROP CALCULATION
          C
000051       30 IF(NART.EQ.0) RETURN
000054          WW=FWDOT
000056          DLAST=0.
000057          IF(NN.NE.1) GO TO 40
000061          SAVW(MARK,KK)=WW
000065          ART=0.
          C
          C     ADDS IN CONTRIBUTION FROM DROP 1 FOR INTERPOLATION WITH LAST DROP
          C
000065          DO 35 I=1,NART
000067       35 ART=ART+FLOAT(I)/FLOAT(NART+1)*WW
000100          FWDOT=FWDOT+ART
000101          RETURN
000102       40 IF(NN.EQ.NP) DLAST=1.
000106          ART=0.
000107          DO 50 I=1,NART
000111       50 ART=ART+SAVW(MARK,KK)+FLOAT(I)/FLOAT(NART+1)*(WW-SAVW(MARK,KK))
                1  +DLAST*(WW-FLOAT(I)/FLOAT(NART+1)*WW)
000136          SAVW(MARK,KK)=FWDOT
000142          FWDOT=FWDOT+ART
000143          RETURN
000144          END
```

```
                SUBROUTINE OUTPUT
000002          COMMON WWW(3000),TTT(3000),W(400),INDEX,KK2,VEL,VD,AKG,DD,PPV,H,DO
               1M1,SIT,S,TIME,J,K,JA,NA,NC,DX,DT,ST,SN,STN,WDOT,P,TF
000002          COMMON SAVW(30,400),MARK,KK,NART,NN,NP
000002          COMMON X(64),Z(64),AB(64),AC(64),AD(64),UO(64),U(64),UN(64),T(64),
               1R(64)
          C
000002          DUM = TIME
000004          WRITE (6,40) K,DUM,SN,ST,WDOT
000021          NH = (NC+1)/2
000024          NM = NC/2
000025          DO 10 N = 1,NH
000027          L = NM+N
000031       10 WRITE (6,30) N,R(N),T(N),L,R(L),T(L)
000053          IF (K.EQ.0) GO TO 20
000054          REY = DOM1**2
000055          DUM = (S/SIT)**3
000060          WRITE (6,50) H,REY,DUM,DT
000073          WRITE(6,70)VEL,VD,AKG,DD,PPV
000111          WRITE (6,60) P,TF
000121       20 CONTINUE
000121          J = 0
000122          RETURN
          C
000123       30 FORMAT (I9,2E15.5(I9,2E15.5)
000123       40 FORMAT (1H0//5X,4HSTEPI6,10X,5HTIME=E12.4,5X,3HRS=E12.4/24X,6HRSDO
               1T=E12.4,5X,5HWDOT=E12.4//18X,1HR,13X,1HT,24X,1HR,13X,1HT)
000123       50 FORMAT (2X,2HH=E10.3,1X,4HREY=E10.3,1X,11HMASS RATIO=E10.3,1X,3HDT
               1=E10.3)
000123       60 FORMAT (2X,2HP=E10.3,1X,3HTF=E10.3)
000123       70 FORMAT(2X,4HVEL=E10.3,3HVD=E10.3,4HAKG=E10.3,3HDD=E10.3,4HPPV=E10.
               13)
000123          END
```

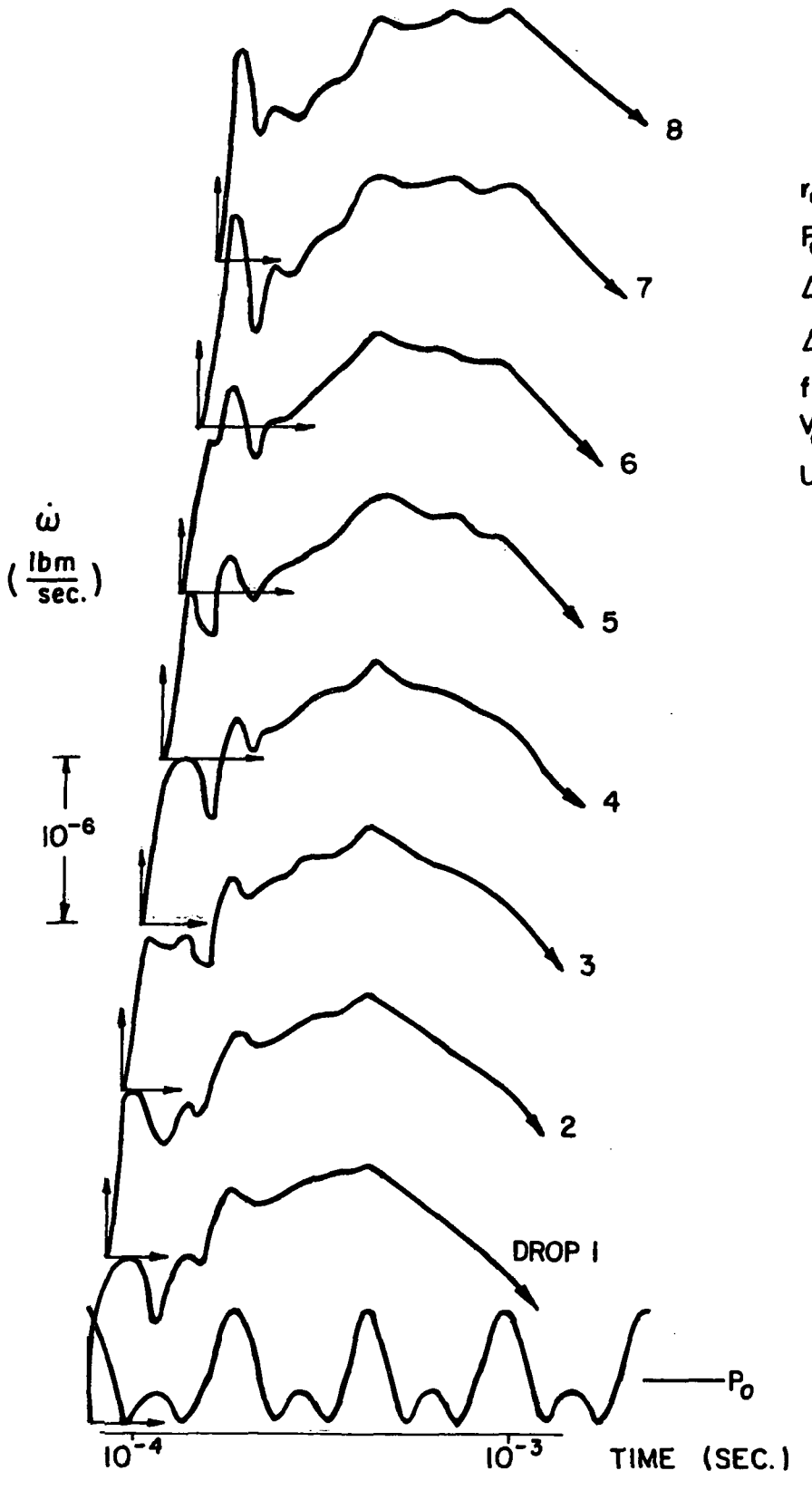
8
7
6
5
4
3
2
DROP 1
$\dot{\omega}$ ($\frac{lbm}{sec.}$)
10^{-6}
P_o
10^{-4}
10^{-3}
TIME (SEC.)
r_o = 50 MICRONS
P_c = 300 psi
$\Delta P_1 = 0.2 P_c$
$\Delta P_2 = 0.8 \Delta P_1$
f = 3000 CPS
V_d = 100 FT./SEC.
U_{gf} = 1200 FT./SEC.

FIG. 1

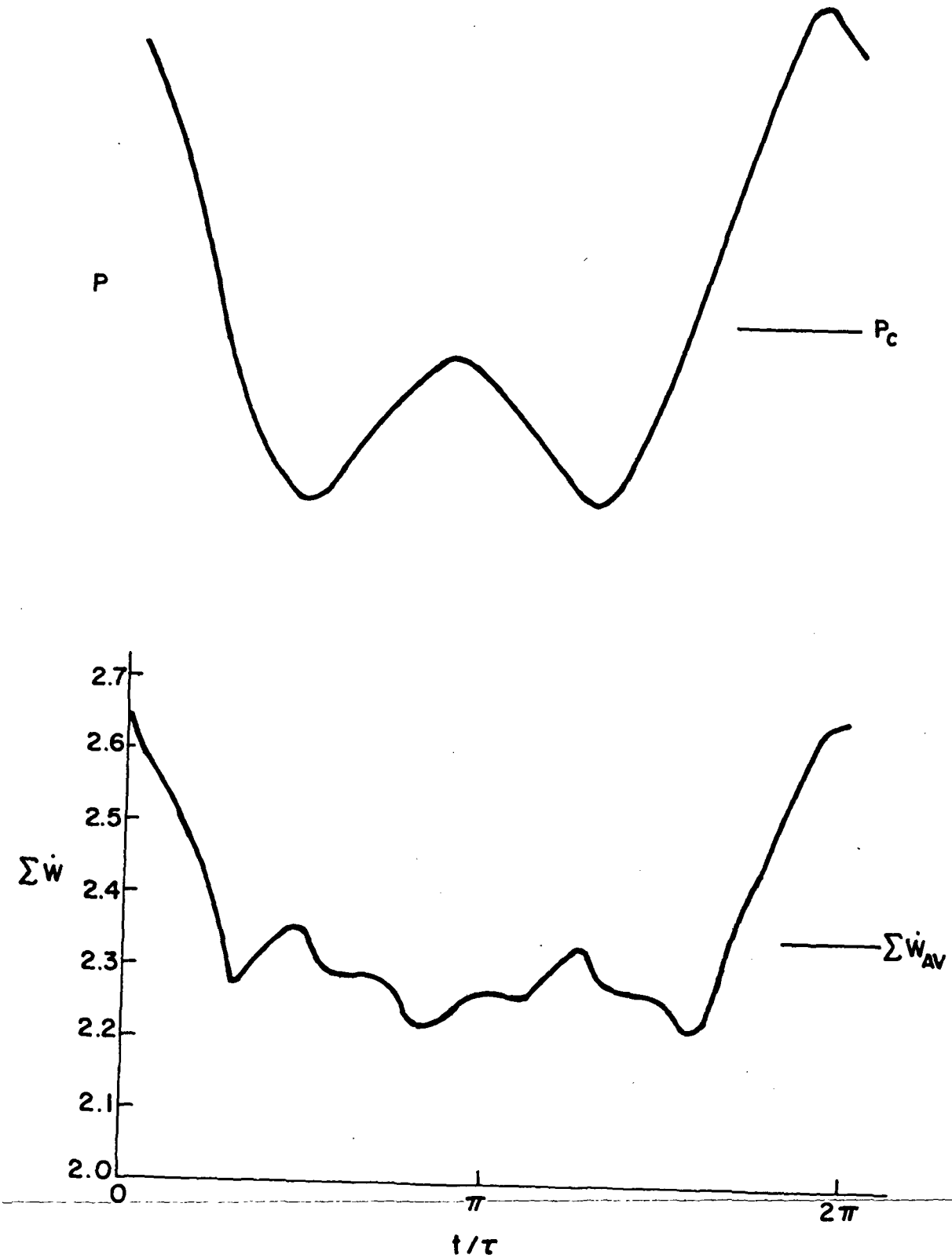

FIG. 2 ARRAY EVAPORATION RATE VS. TIME

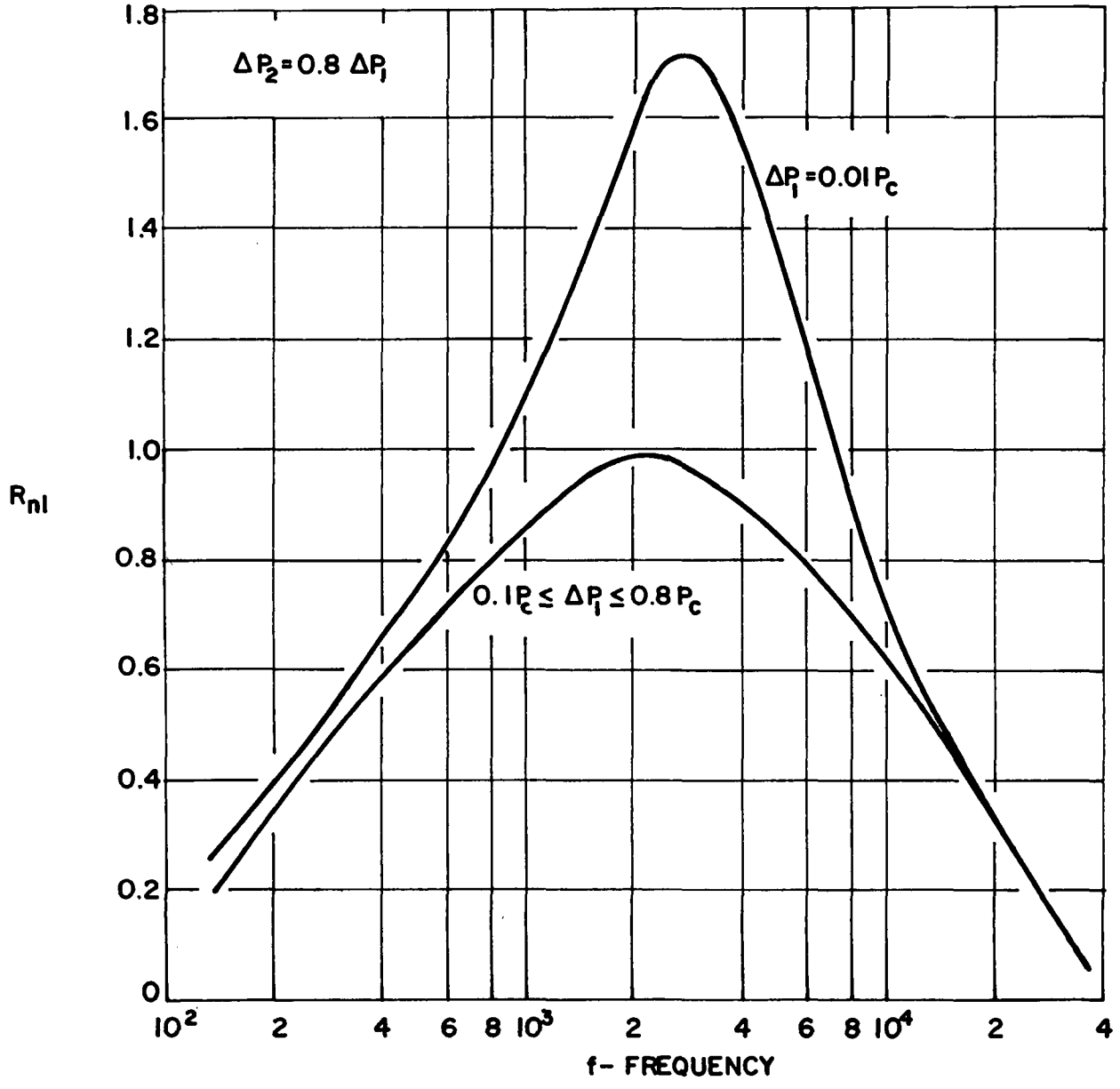

FIG. 3 RESPONSE FACTOR VS. FREQUENCY: EFFECT OF AMPLITUDE

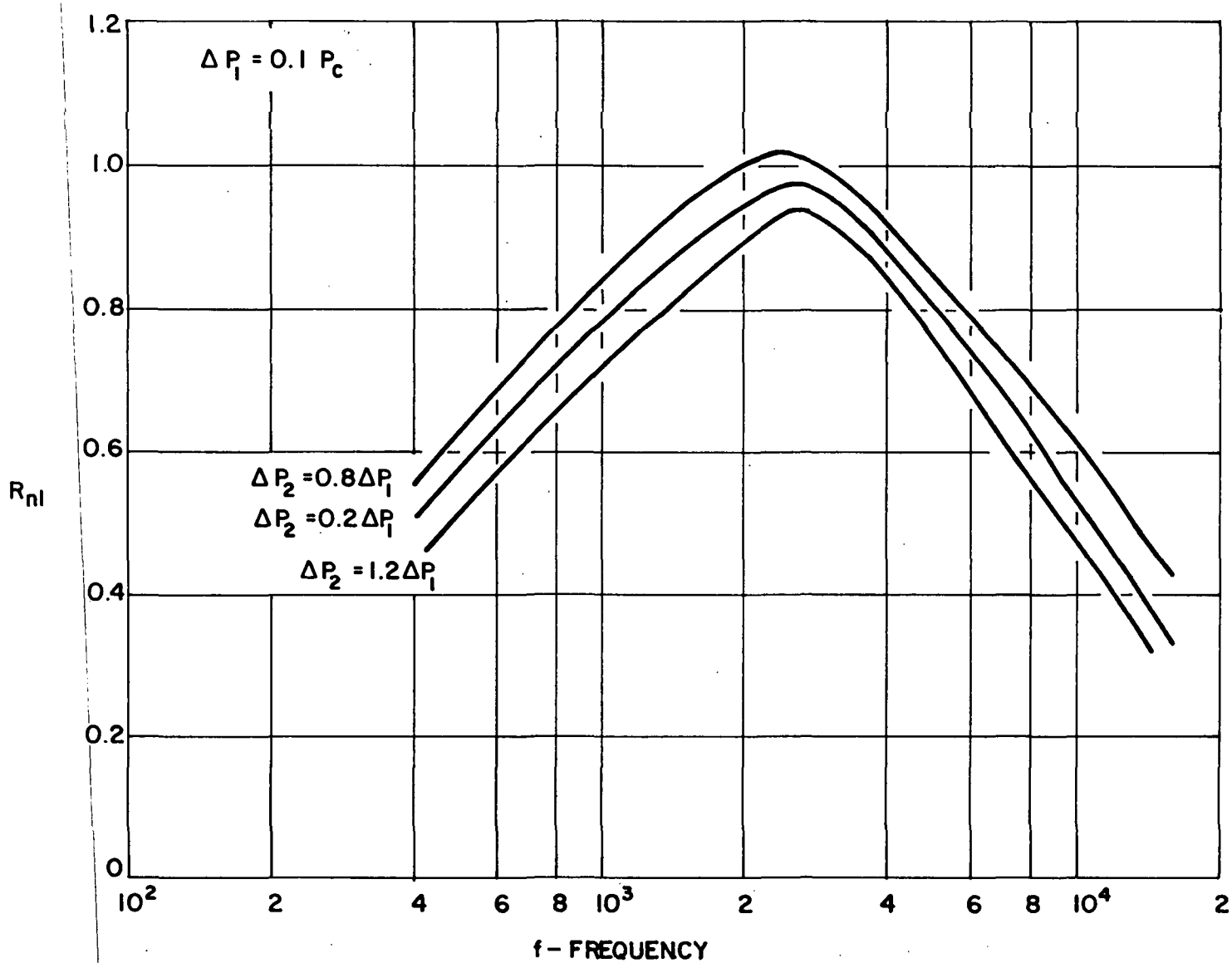

FIG. 4a RESPONSE FACTOR VS. FREQUENCY: EFFECT OF DISTORTION $\Delta P_1 = 0.1\ P_c$

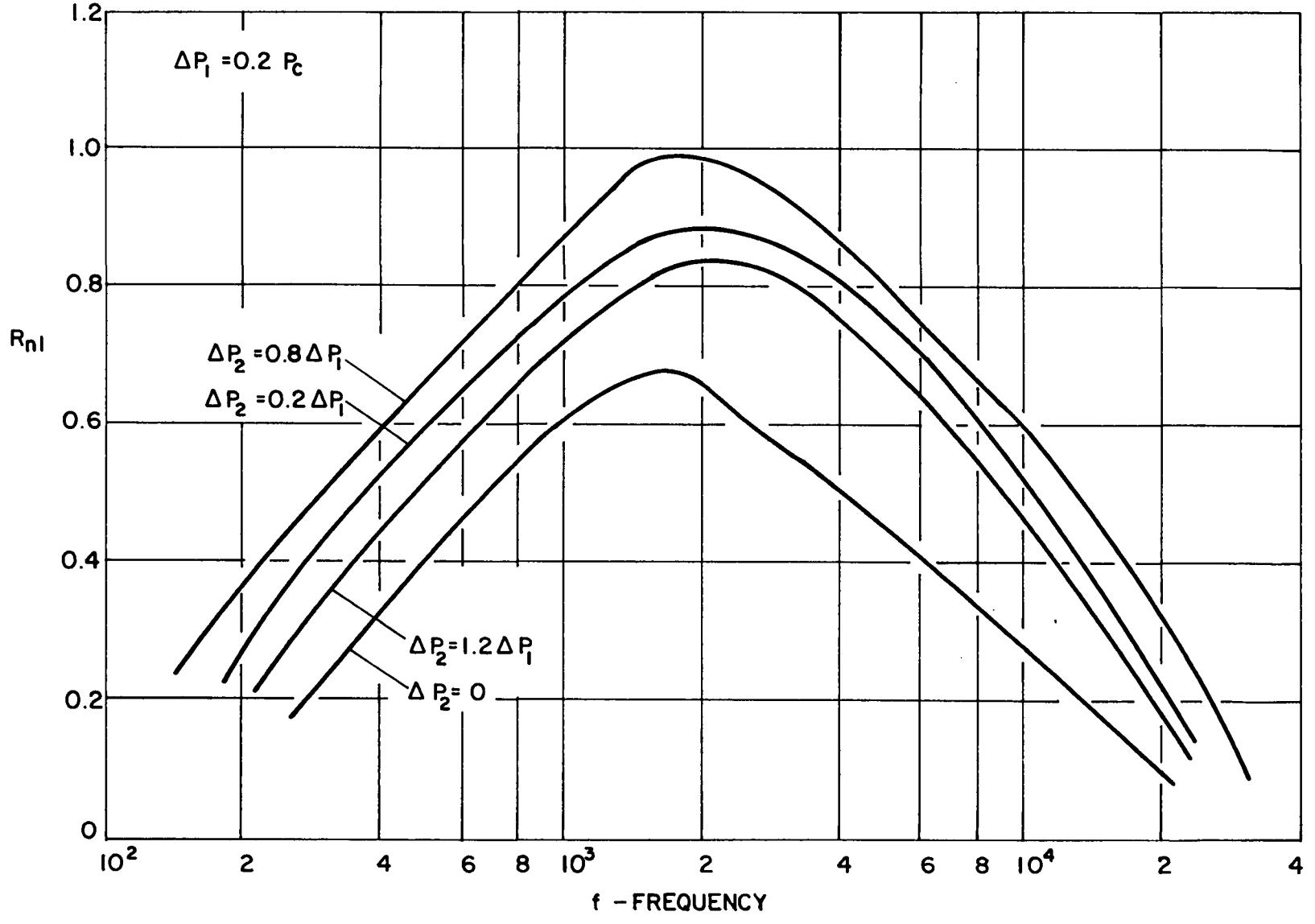

FIG. 4b RESPONSE FACTOR VS. FREQUENCY: EFFECT OF DISTORTION
$\Delta P_1 = 0.2\ P_c$

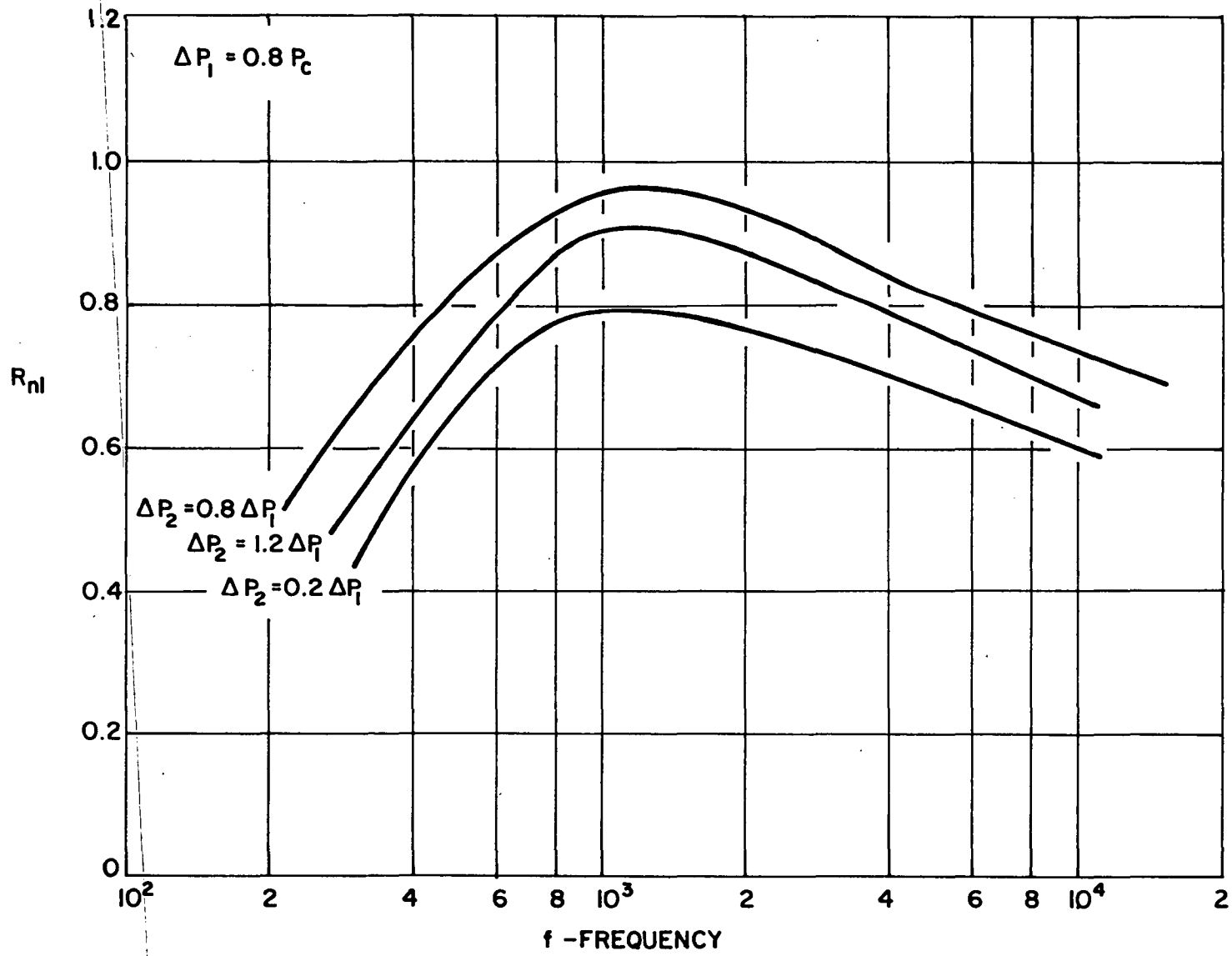

FIG. 4c RESPONSE FACTOR VS. FREQUENCY: EFFECT OF DISTORTION $\Delta P_1 = 0.8\ P_c$

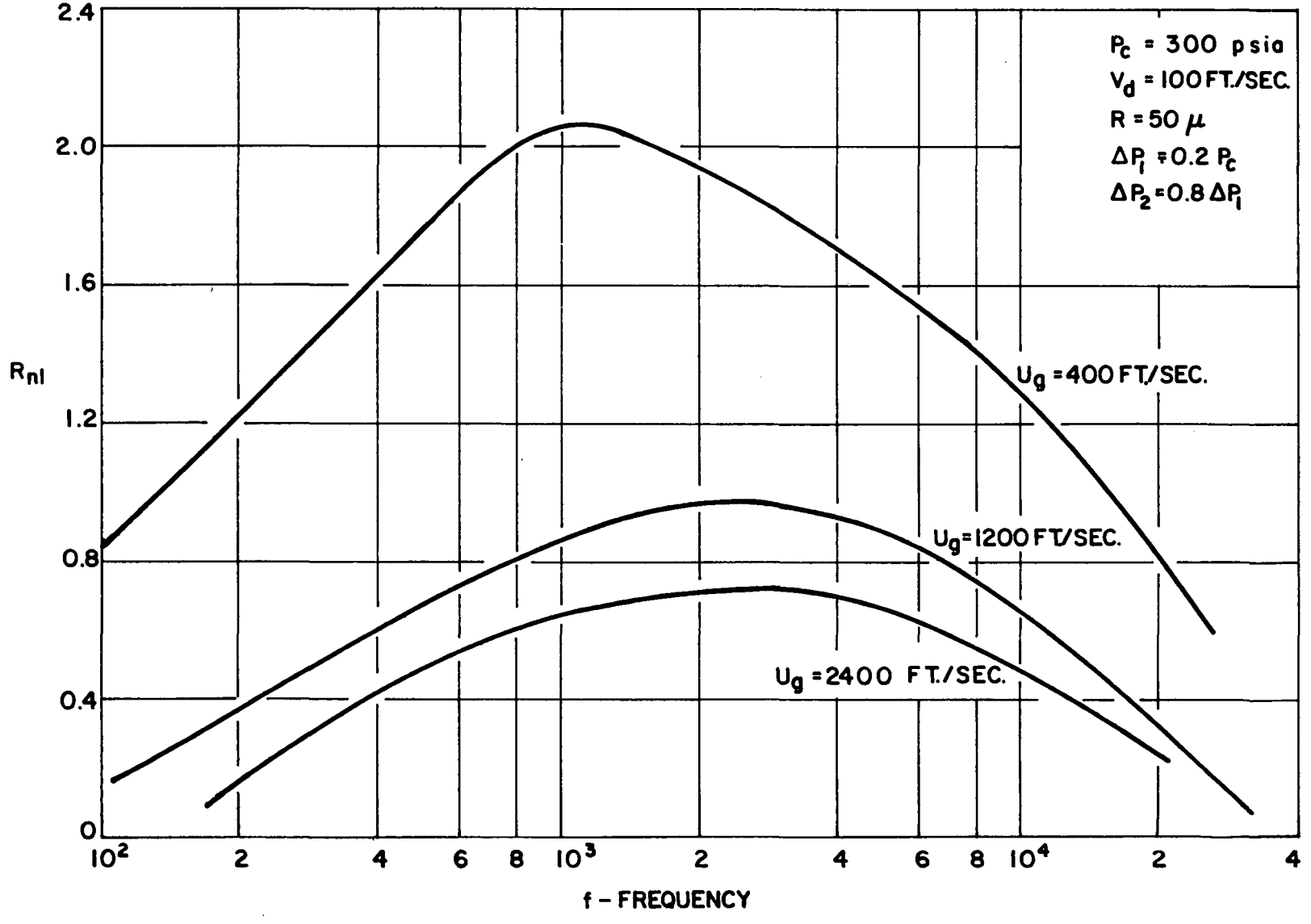

FIG. 5 RESPONSE FACTOR VS. FREQUENCY: EFFECT OF FINAL GAS VELOCITY

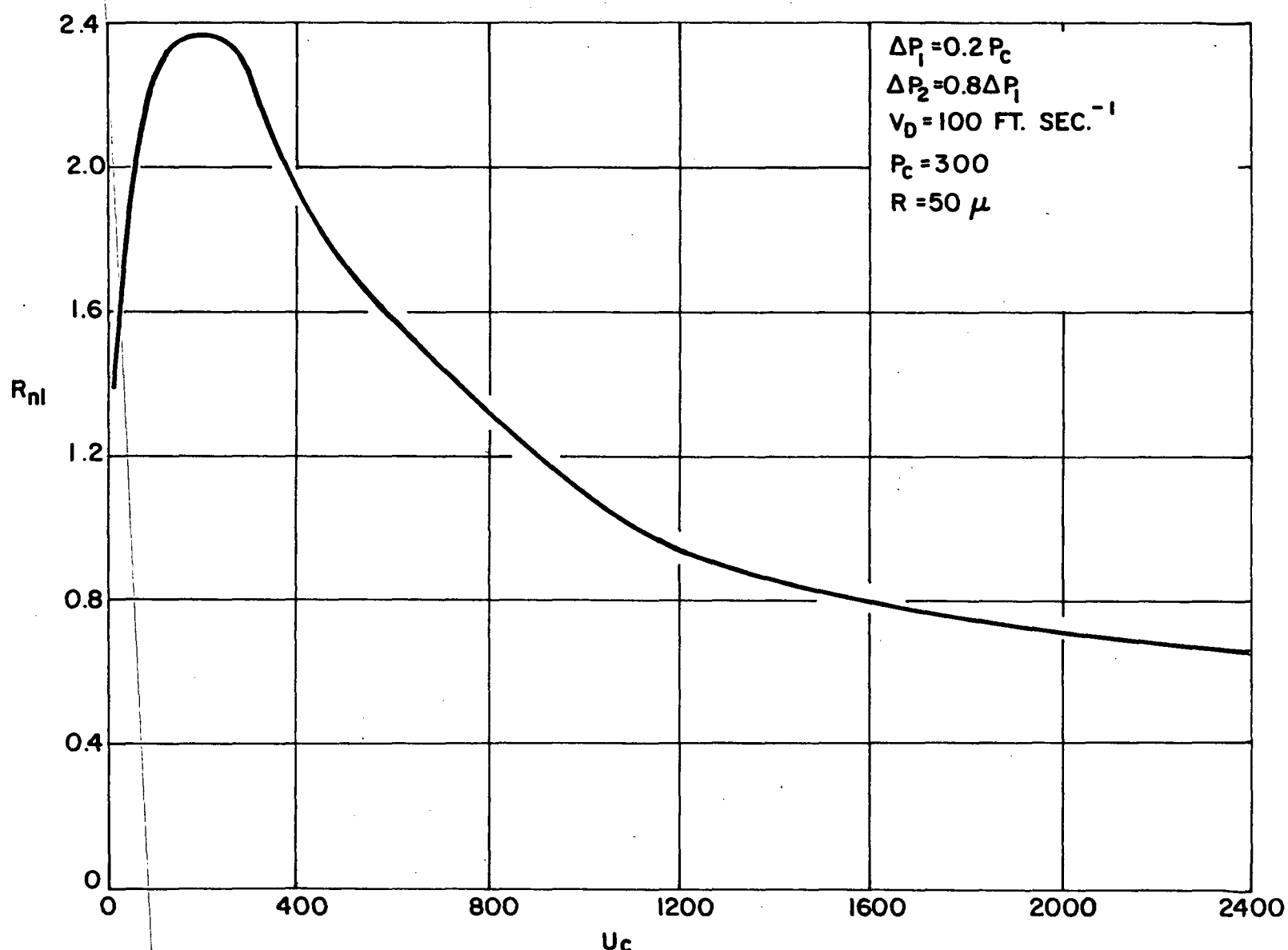

FIG. 6 RESPONSE FACTOR VS. FINAL GAS VELOCITY

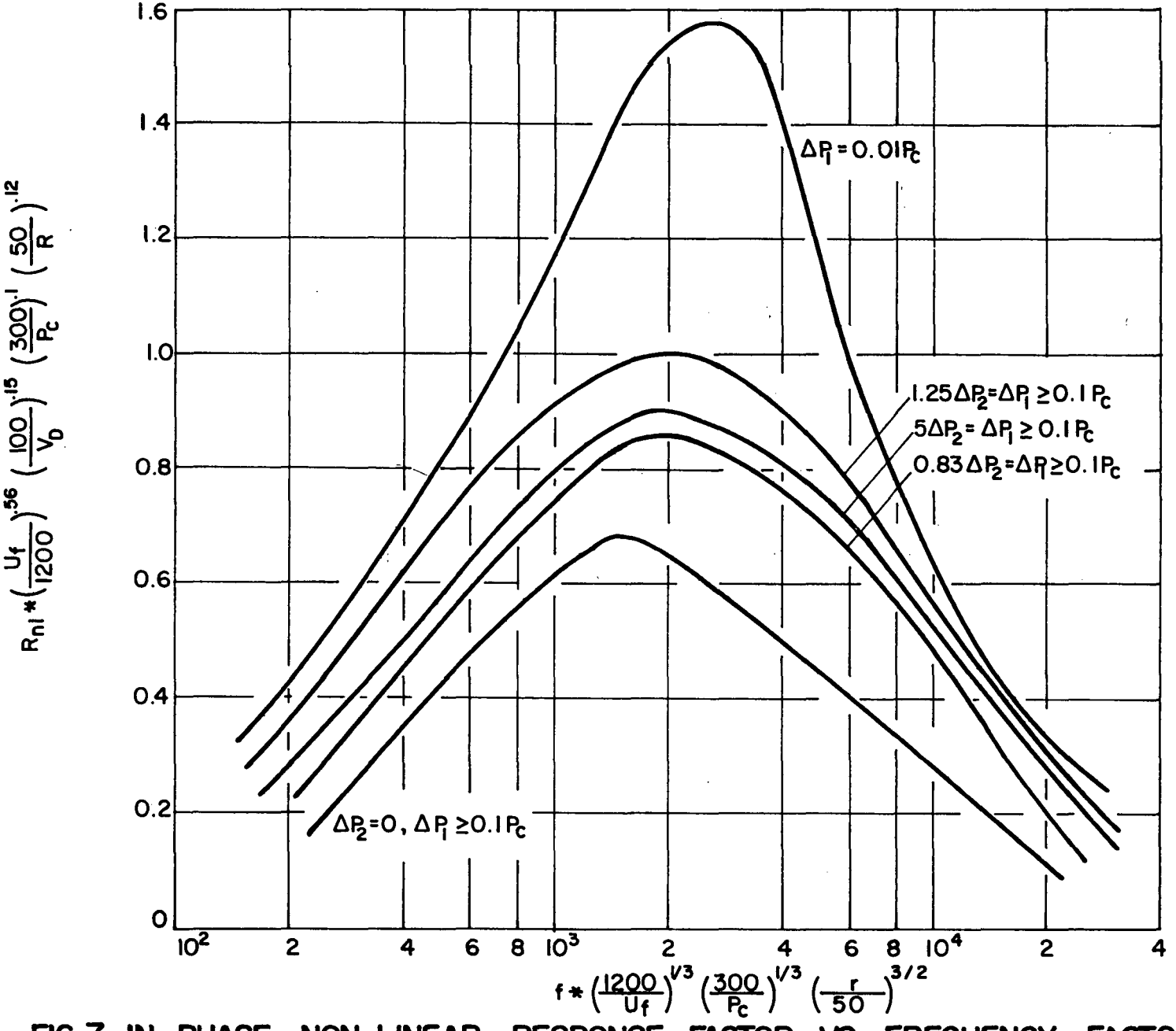

FIG.7 IN PHASE NON-LINEAR RESPONSE FACTOR VS. FREQUENCY FACTOR

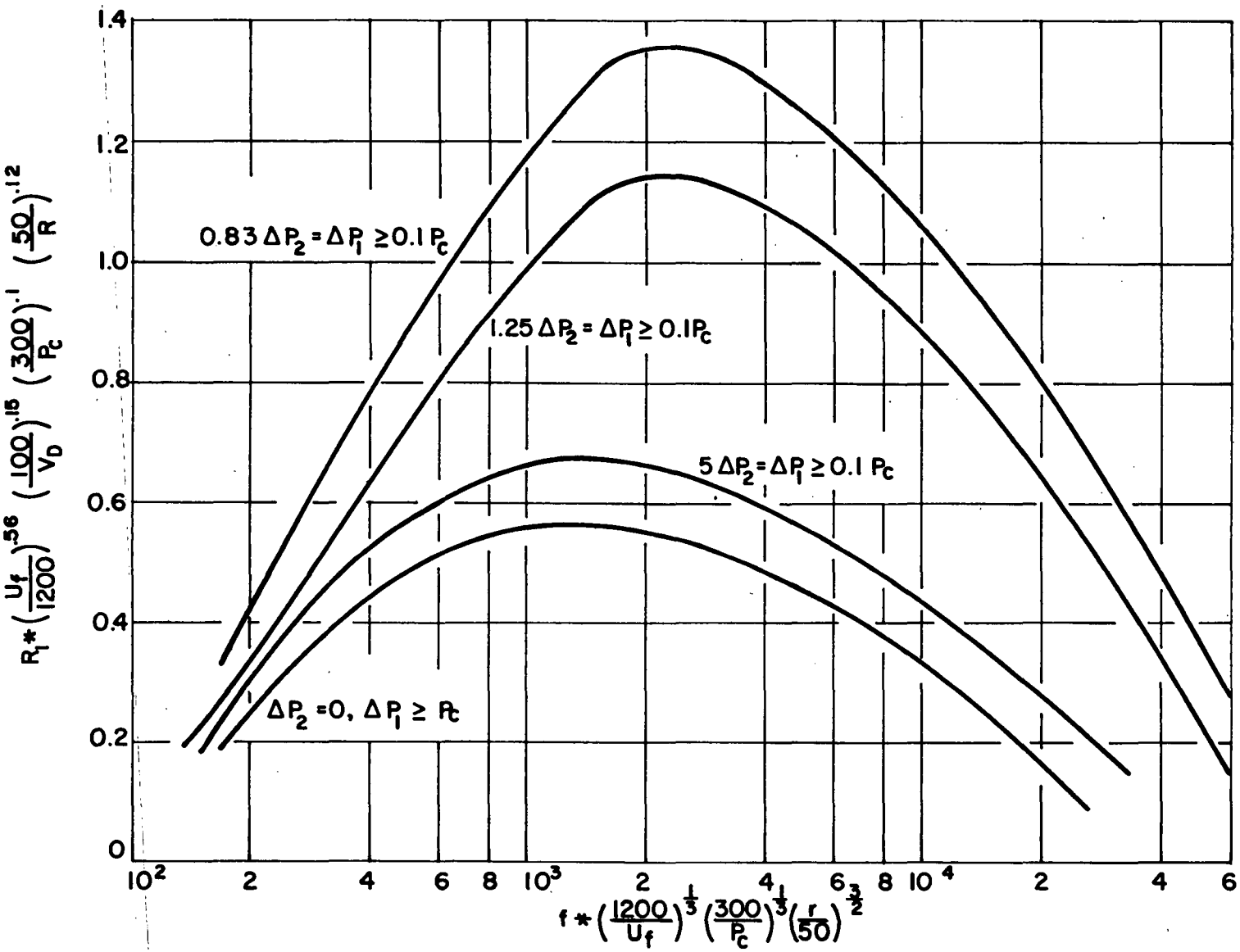

FIG. 8 IN PHASE FUNDAMENTAL RESPONSE FACTOR VS. FREQUENCY FACTOR

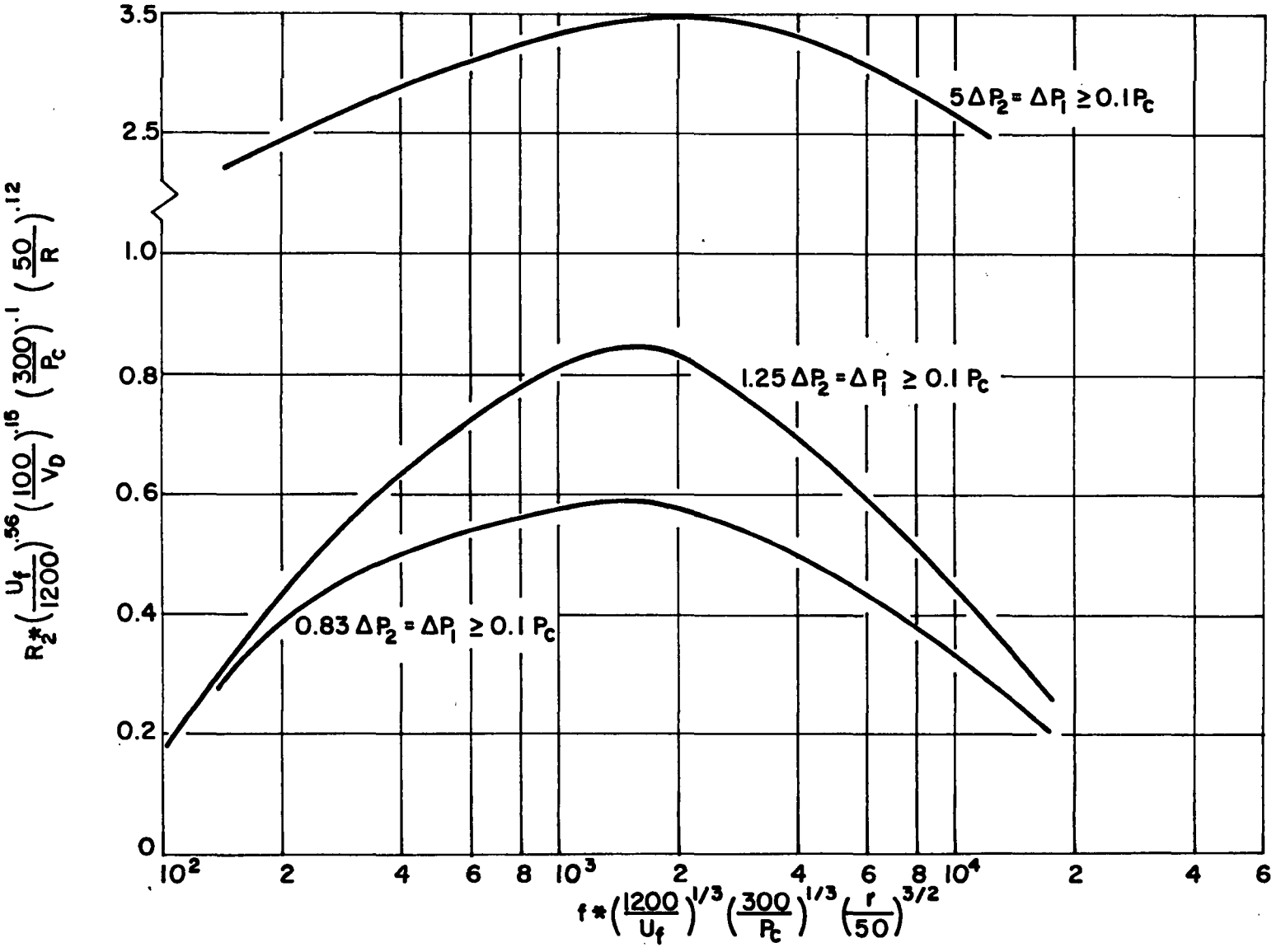

FIG.9 FIRST HARMONIC RESPONSE FACTOR VS. FREQUENCY FACTOR

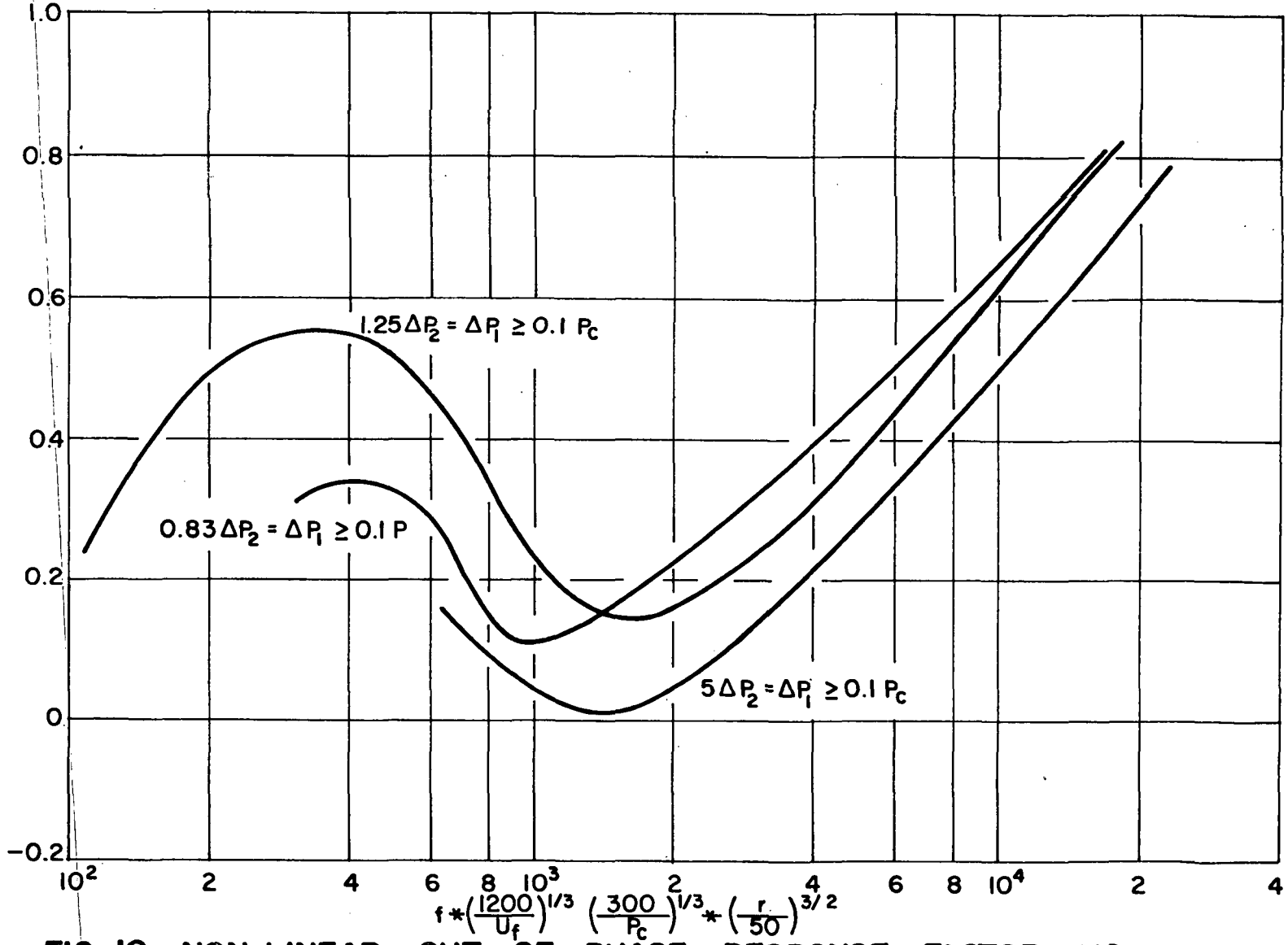

FIG. 10 NON-LINEAR OUT OF PHASE RESPONSE FACTOR VS. FREQUENCY FACTOR

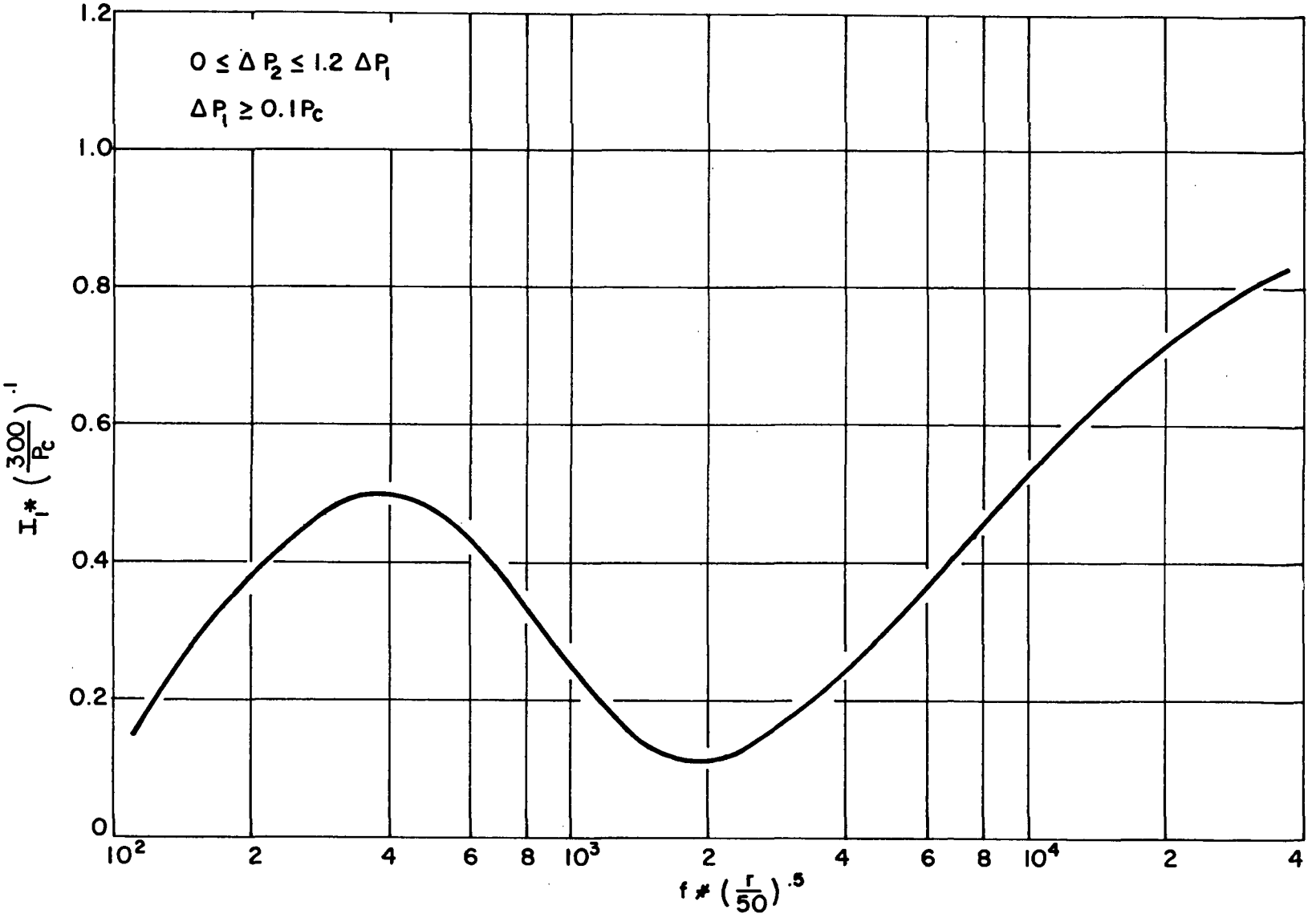

FIG. 11 FUNDAMENTAL OUT OF PHASE RESPONSE FACTOR VS. FREQUENCY FACTOR

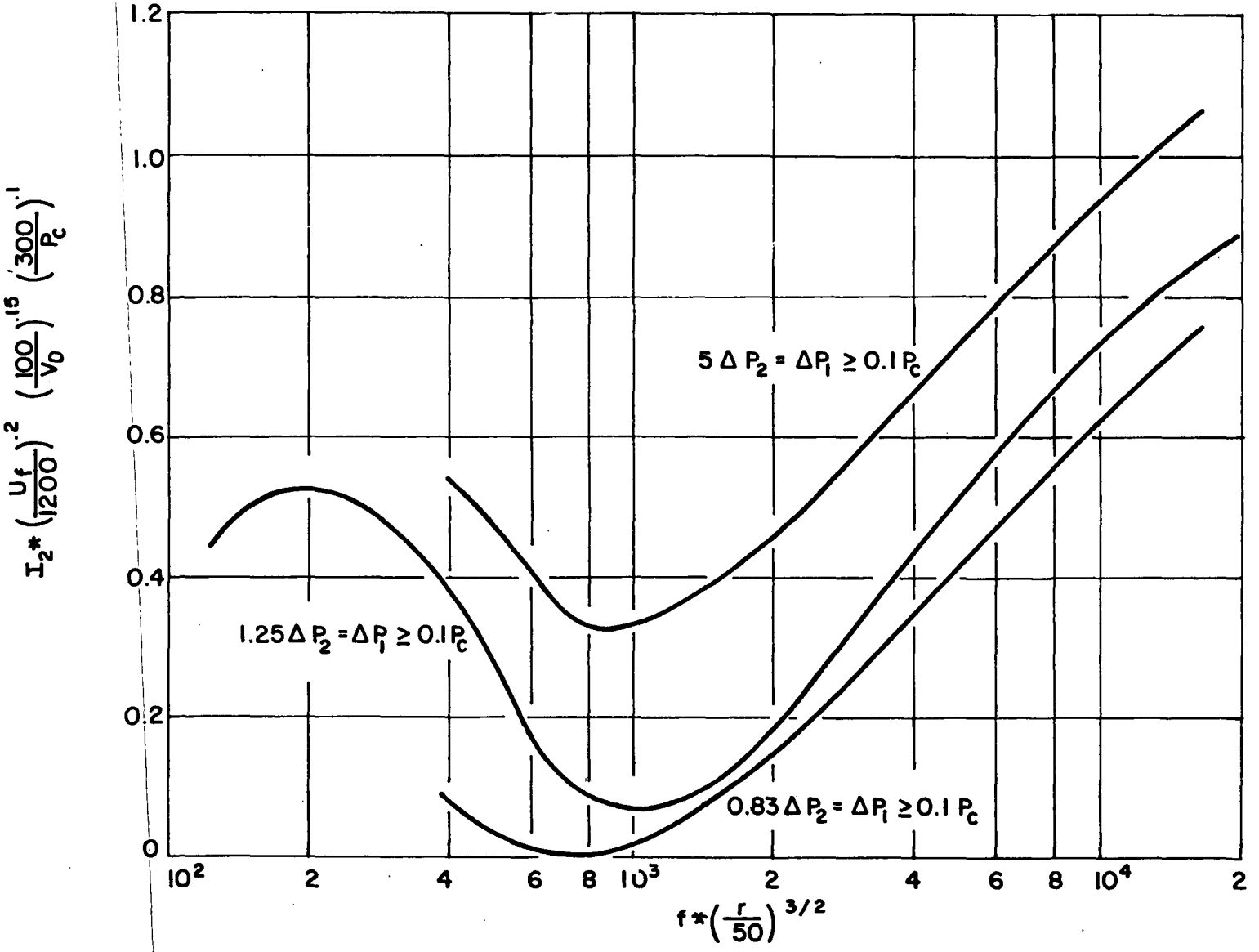

FIG. 12 FIRST HARMONIC OUT OF PHASE RESPONSE FACTOR VS. FREQUENCY FACTOR

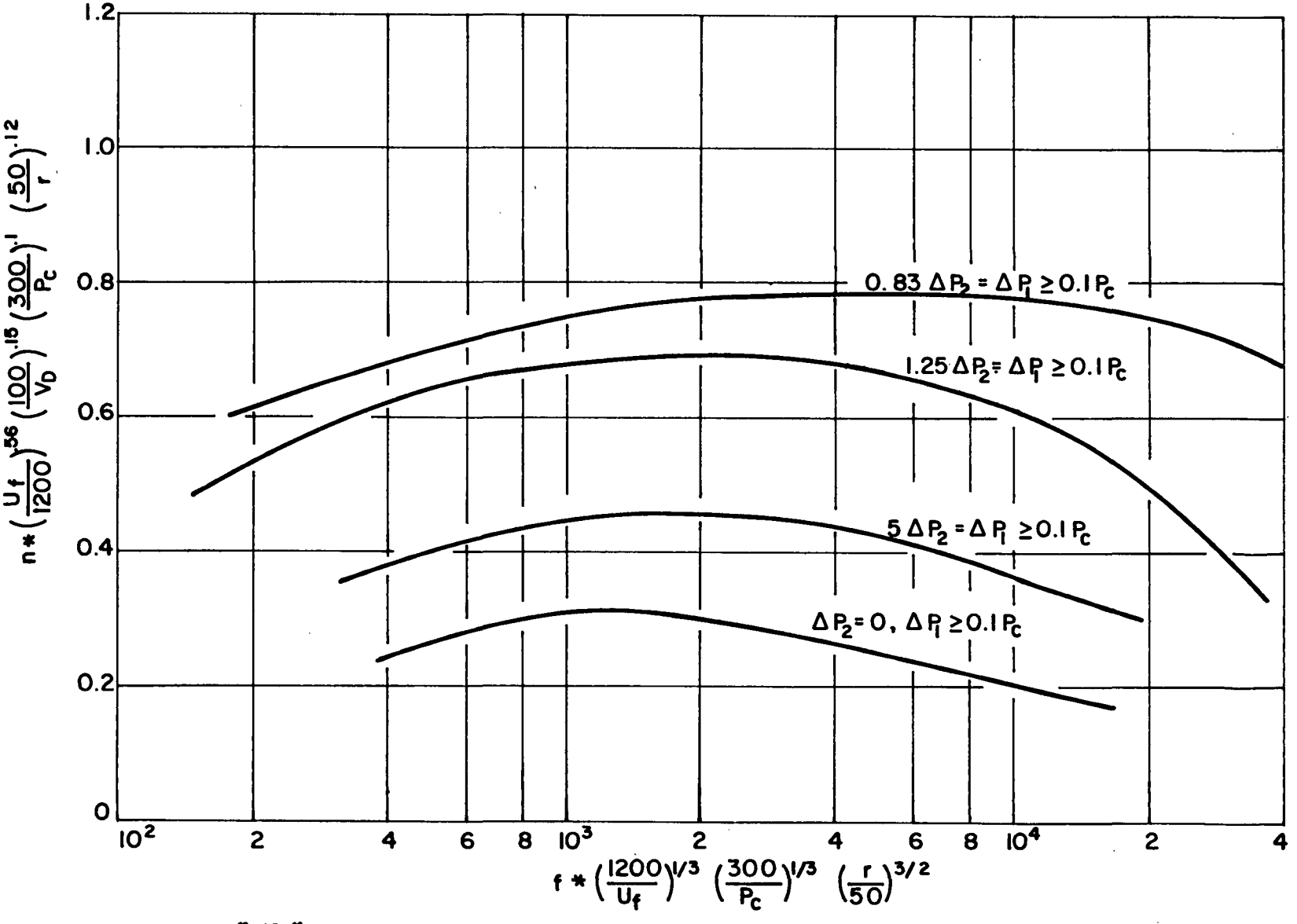

FIG. 13 "n" VS. FREQUENCY FACTOR

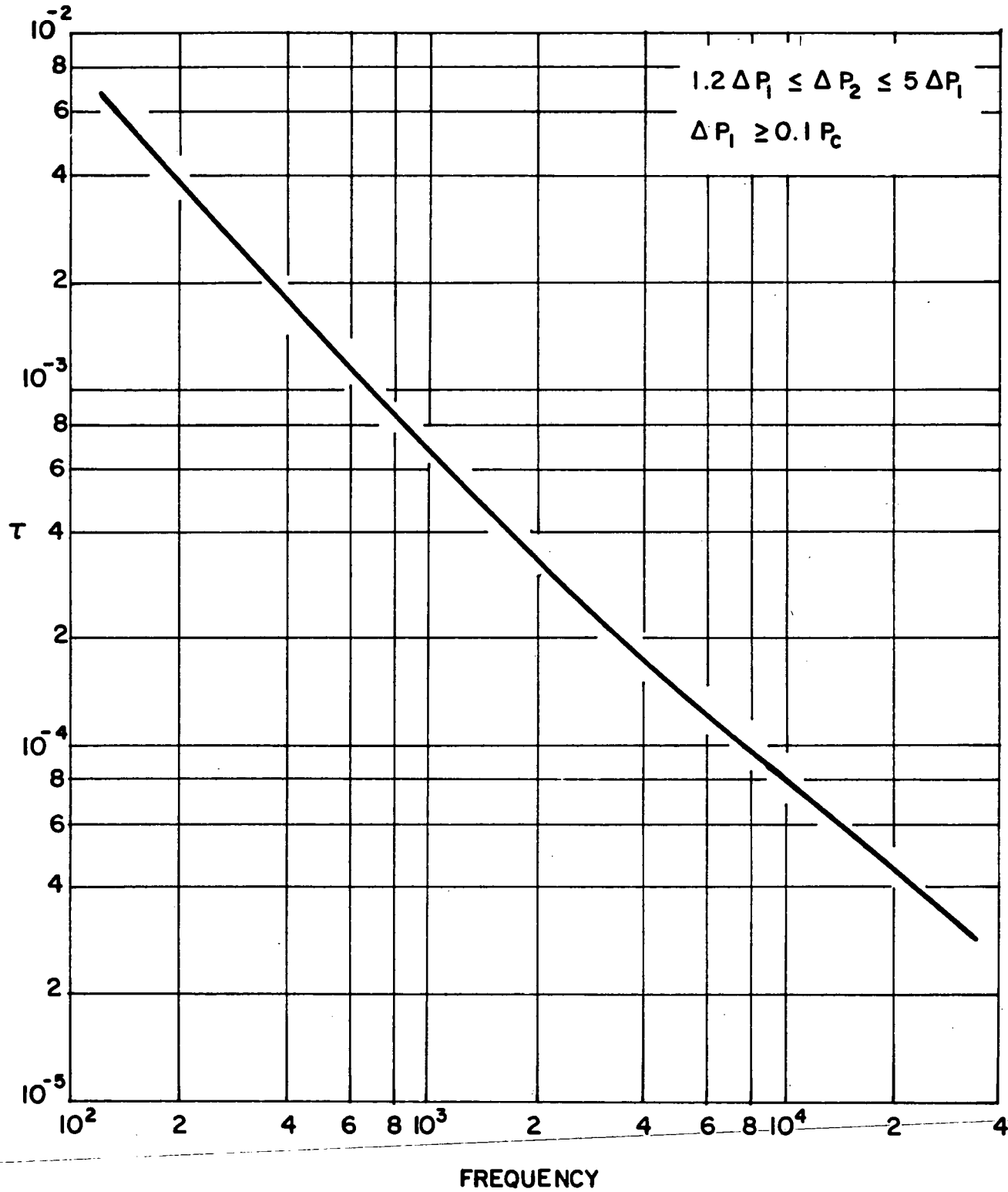

FIG. 14 τ VS. FREQUENCY

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